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(57) Abstract

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HIV ENVELOPE POLYPEPTIDES AND VACCINE

BACKGROUND OF THE INVENTION5 Field of the Invention

This invention relates to HIV envelope polypeptides and vaccines containing the polypeptides.

Description of the Related Art

10 Acquired immunodeficiency syndrome (AIDS) is caused by a retrovirus identified as the human immunodeficiency virus (HIV). There have been intense efforts to develop a vaccine that induces a protective immune response based on induction of antibodies or
15 cellular responses. Recent efforts have used subunit vaccines where an HIV protein, rather than attenuated or killed virus, is used as the immunogen in the vaccine for safety reasons. Subunit vaccines generally include gp120, the portion of the HIV envelope protein
20 which is on the surface of the virus.

The HIV envelope protein has been extensively described, and the amino acid and nucleic acid sequences encoding HIV envelope from a number of HIV strains are known (Myers, G. et al., 1992. Human
25 Retroviruses and AIDS. A compilation and analysis of nucleic acid and amino acid sequences. Los Alamos National Laboratory, Los Alamos, New Mexico). The HIV envelope protein is a glycoprotein of about 160 kd (gp160) which is anchored in the membrane bilayer at
30 its carboxyl terminal region. The N-terminal segment, gp120, protrudes into the aqueous environment surrounding the virion and the C-terminal segment, gp41, spans the membrane. Via a host-cell mediated process, gp160 is cleaved to form gp120 and the
35 integral membrane protein gp41. As there is no covalent attachment between gp120 and gp41, free gp120 is sometimes released from the surface of virions and

infected cells.

The gp120 molecule consists of a polypeptide core of 60,000 daltons which is extensively modified by N-linked glycosylation to increase the apparent molecular weight of the molecule to 120,000 daltons. The amino acid sequence of gp120 contains five relatively conserved domains interspersed with five hypervariable domains. The positions of the 18 cysteine residues in the gp120 primary sequence, and the positions of 13 of the approximately 24 N-linked glycosylation sites in the gp120 sequence are common to all gp120 sequences. The hypervariable domains contain extensive amino acid substitutions, insertions and deletions. Sequence variations in these domains result in up to 30% overall sequence variability between gp120 molecules from the various viral isolates. Despite this variation, all gp120 sequences preserve the ability of the virus to bind to the viral receptor CD4 and to interact with gp41 to induce fusion of the viral and host cell membranes.

gp120 has been the object of intensive investigation as a vaccine candidate for subunit vaccines, as the viral protein which is most likely to be accessible to immune attack. At present, clinical trials using gp120 MN strain are underway. However, to date no human vaccine trial has been of sufficient size to confirm or refute vaccine efficacy.

The development of candidate HIV-1 vaccines is burdened by the lack of in vivo or in vitro models of HIV-1 infection that accurately approximate the conditions of natural infection in humans. Several candidate HIV-1 vaccines [Berman et al.; *J. Virol.* 7:4464-9 (1992); Haigwood et al.; *J. Virol.* 66:172-82 (1992); Salmon-Ceron et al.; *AIDS Res. and Human Retroviruses* 11:1479-86 (1995)] have been described that elicit broadly cross-reactive antibodies able to

neutralize a variety of diverse HIV-1 isolates in vitro. However, the relevance of in vitro assays to protective immunity in vivo is uncertain. Although several vaccines have provided chimpanzees with protection from challenge by homologous and heterologous strains of HIV-1, protection has not always correlated with in vitro neutralization assays carried out in T cell lines, or in lectin- and cytokine-activated peripheral blood mononuclear cells (PBMCs) [Berman et al.; *Nature* 345:622-5 (1990); Bruck et al.; *Vaccine* 12(12):1141-8 (1994); El-Amad et al.; *AIDS* 9:1313-22 (1995); Girard et al.; *J. Virol.* 69:6239-48 (1995); and Fulz et al.; *Science* 256:1687-1690 (1992)]. While successful protection of chimpanzees is encouraging and has historically proved to be a reliable indicator of vaccine efficacy, the conditions of infection in all experimental models of HIV-1 infection differ significantly from natural infection in humans.

Experimental HIV-1 infection in vivo and in vitro both suffer from the limitation that the in vitro amplification of HIV-1, which is required to prepare virus stocks for in vitro or in vivo infectivity experiments, imposes a genetic selection that results in a spectrum of virus quasi-species that differ from the spectrum of variants present in the clinical specimens used to establish the culture [Kusumi et al.; *J. Virol.* 66:875 (1992); Meyerhans et al.; *Cell* 58:901-10 (1989)]. Because of these uncertainties, and even greater uncertainties related to the amount of virus transmitted, the site and cell type involved in initial replication, and the kinetics of virus dissemination, the ability of currently available in vitro or in vivo assays to reliably predict vaccine efficacy is questionable.

One of the candidate HIV-1 vaccines that have

entered human clinical trials is recombinant gp120 prepared in Chinese hamster ovary (CHO) cells from the MN strain of HIV-1 (MN-rgp120) (Berman et al.; *J. Virol.* 7:4464-9 (1992)). To date, approximately 499 adults have participated in Phase 1 and 2 immunogenicity and safety trials of this vaccine. The data collected thus far suggest that MN-rgp120 is safe, immunogenic, and elicits high titers of neutralizing antibodies in greater than 95% of individuals immunized according to a 0, 1, and 6 month immunization schedule [Belshe et al.; *JAMA* 272(6):475-80 (1994); McElrath; *Seminars in Cancer Biol.* 6:1-11 (1995)]. However, during the course of these trials, nine vaccinees who received MN-rgp120 have become infected with HIV-1 through high risk behavior. Small trials, such as these, in populations with low rates of infection and minimally sized placebo control groups do not have sufficient statistical power to confirm or refute vaccine efficacy.

However, effective vaccines based on gp120 or another HIV protein for protection against additional strains of HIV are still being sought to prevent the spread of this disease.

Description of the Background Art

Recombinant subunit vaccines are described in Berman et al., PCT/US91/02250 (published as number WO91/15238 on 17 October 1991). See also, e.g. Hu et al., *Nature* 328:721-724 (1987) (vaccinia virus-HIV envelope recombinant vaccine); Arthur et al., *J. Virol.* 63(12): 5046-5053 (1989) (purified gp120); and Berman et al., *Proc. Natl. Acad. Sci. USA* 85:5200-5204 (1988) (recombinant envelope glycoprotein gp120).

Numerous sequences for gp120 are known. The sequence of gp120 from the IIIB substrain of HIV-1_{LAI}

referred to herein is that determined by Muesing et al., "Nucleic acid structure and expression of the human AIDS/lymphadenopathy retrovirus, *Nature* 313:450-458 (1985). The sequences of gp120 from the NY-5, Jrscf, 26, Z321, and HXB2 strains of HIV-1 are listed by Myers et al., "Human Retroviruses and AIDS; A compilation and analysis of nucleic acid and amino acid sequences," Los Alamos National Laboratory, Los Alamos, New Mexico (1992). The sequence of the Thai isolate A244 is provided by McCutchan et al., "Genetic Variants of HIV-1 in Thailand," *AIDS Res. and Human Retroviruses* 8:1887-1895 (1992). The MN₁₉₈₄ clone is described by Gurgo et al., "Envelope sequences of two new United States HIV-1 isolates," *Virology* 164: 531-536 (1988). As used herein, MN, MN-rgp120, the MN clone or isolate refers to MN_{GNE}. The MN_{GNE} amino acid sequence is Sequence ID No. 29.

Each of the above-described references is incorporated herein by reference in its entirety.

Summary of the Invention

Oligonucleotide sequences encoding gp120 polypeptides from breakthrough isolates of vaccine trials using MN-rgp120 and the encoded gp120 polypeptides are provided. Use of the gp120 polypeptides from one or more of the isolates in a subunit vaccine, usually together with MN-rgp120, can provide protection against HIV strains that are sufficiently different from the vaccine strain (e.g.; MN-rgp120) that the vaccine does not confer protection against those strains. Antibodies induced by the polypeptides are also provided.

Brief Description of the Drawings

Figure 1 illustrates the kinetics of antibody response to MN-rgp120 in vaccinees infected with HIV-1.

Sera were collected at the time points indicated and assayed for antibodies reactive with MN-rgp120 (open circles) or a synthetic peptide derived from the V3 domain of MN-rgp120 (closed circles). Arrows indicate dates of injection. Plus sign indicates the first time HIV-1 infection was detected. Shaded area indicates data collected after HIV-1 infection. Data from vaccinee C6 is shown in panel A; C8 in panel B; C7, panel C; C11, panel D; C10, panel E; C17, panel F; and C15, panel G.

Figure 2 illustrates the kinetics of CD4 blocking antibody response in vaccinees infected with HIV-1. Sera were collected at the time points indicated and assayed for antibodies able to block the binding of [¹²⁵I]-labeled MN-rgp120 to cell surface CD4. Arrows indicate dates of injection. Plus sign indicates the first time HIV-1 infection was detected. Shaded area indicates data collected after HIV-1 infection. Data from vaccinee C6 is shown in panel A; C8 in panel B; C7, panel C; C11, panel D; C10, panel E; C17, panel F; and C15, panel G.

Figure 3 illustrated predicted amino acid sequences of envelope glycoproteins (gp120) from breakthrough viruses. Proviral DNA sequences were amplified by PCR from PBMCs and cloned into the PRK5 expression plasmid. Two clones from each infected vaccinee were sequenced from double stranded plasmid DNA. Sequence numbering is with reference to the initiator methionine residue of gp120. For the purpose of comparison, the sequences shown begin at amino acid 12 of the mature, fully processed, envelope glycoproteins (corresponding to position 41 of the gp120 open reading frame). Shaded areas indicate sequences at neutralizing epitopes, dark boxes indicate polymorphisms thought to be important for the binding of virus neutralizing MABs reactive with MN-rgp120.

Conserved (C) regions and variable (V) regions are indicated above the sequences. Boxes indicate sequence homologies and polymorphisms.

Figure 4 illustrates immunoprecipitation of recombinant gp120 prepared from breakthrough viruses. Recombinant gp120s from the seven breakthrough viruses were prepared by transient transfection of 293s cells. Cells were metabolically labeled with ³⁵S methionine and growth conditioned cell culture supernatants were immunoprecipitated with polyclonal antisera to MN-rgp120. Immunoprecipitates were resolved by SDS-PAGE and visualized by autoradiography. C8 lanes a and b correspond to clones C8.3 and C8.6; C6 lanes a and b correspond to clones C6.1 and C6.5; C7 lanes a and b correspond to clones C7.2 and C7.10; C17 lanes a and b correspond to clones C17.1 and C17.3; C11 lanes a and b correspond to clones C11.5 and C11.7; C10 lanes a and b correspond to clones C10.5 and C10.7; C15 lanes a and b correspond to clones C15.2 and C15.3.

Figure 5 illustrates binding of monoclonal antibodies to recombinant gp120 from breakthrough viruses. Growth-conditioned cell culture supernatants were collected from 293s cells transiently transfected with plasmids directing the expression of breakthrough virus envelope glycoproteins. The relative rgp120 concentrations were determined by ELISA using MAb 5B6 specific for the HSV-1 glycoprotein D flag epitope at the amino terminus of all of the rgp120 variants described herein. The resulting rgp120 preparations were captured onto wells of microtiter plates coated with a polyclonal antibody specific for a conserved sequence in the C-terminus of gp120. The binding of virus neutralizing monoclonal antibodies reactive with gp120 was determined by ELISA. A, binding by MAb (5B6) specific for the HSV-1 glycoprotein D flag epitope; B, binding by MAb (1034) against the V3 domain of

MN-rgp120; C binding by MAb (50.1) raised against a synthetic peptide corresponding to the V3 domain of MN-rgp120; D, binding by a human MAb (15e) known to block the binding of gp120 to CD4.

5 Figure 6 depicts the mature envelope glycoprotein (gp120) from the MN clone of the MN strain of HIV-1 (SEQ. ID. NO. 29). Hypervariable domains are indicated in bold, and the V and C regions are indicated (according to Modrow et al., J. Virology 61(2):570
10 (1987). Potential glycosylation sites are marked with a (*).

Detailed Description of the Invention

15 The present invention provides gp120 polypeptides from breakthrough isolates of HIV vaccine trials. Novel oligonucleotide sequences encoding gp120 from breakthrough isolates which can be used to express gp120 are also provided. Use of gp120 polypeptides from one or more of the isolates in a subunit vaccine,
20 usually together with MN-rgp120, can provide protection against HIV strains that are sufficiently different from the vaccine strain (e.g.; MN-rgp120) that the vaccine does not confer protection against those strains.

25 In one embodiment, the vaccine is based on the use of the MN-rgp120 polypeptide (Sequence ID No. 29) and gp120 polypeptides from MN-like viruses that include neutralizing epitopes that are not present in the initial vaccine strain, and are sufficiently different
30 from those of the vaccine strain, to have been able to cause HIV-1 infections in MN-rgp120 vaccinated individuals (i.e.; to result in breakthrough infections). Use of the initial vaccine strain empirically determines the viruses present in the
35 population that contain additional neutralizing epitopes sufficiently different from those of the

vaccine strain to escape protection induced by the vaccine strain. Use of an initial representative gp120 polypeptide in a vaccine acts as a sieve so that viruses that are not effectively protected against by the vaccine strain breakthrough the vaccine, empirically resulting in determination of additional strains in a given geographic region that are not protected against by the initial vaccine strain. Use of gp120 from those breakthrough isolates complements the vaccine isolate by providing additional neutralizing epitopes not present in the initial vaccine strain, therefore creating a more complete vaccine that confers protection against multiple different virus strains in the region.

Prior HIV-1 vaccine strategies were based on selection of appropriate candidate vaccine polypeptides based on homology alignment studies. However, since some of the neutralizing epitopes are conformation-dependent and the location of all of these epitopes is not known, this approach necessarily cannot determine all of the neutralizing epitopes that should be included in a vaccine for a particular region. In contrast, the present approach uses a selected representative strain and empirically determines strains that are sufficiently different and therefore breakthrough the barrier of protection provided by the initial vaccination program. Those strains can be included in the vaccine to confer more complete protection from HIV strains in the region. In addition, those strains can be used alone to confer protection against the breakthrough virus.

In another embodiment, the invention comprises a vaccine containing a first HIV gp120 polypeptide sequence and a breakthrough isolate HIV gp120 polypeptide sequence from a vaccinee vaccinated with a vaccine including the first HIV gp120 polypeptide

sequence, the HIV gp120 polypeptide sequences being in a suitable carrier. Fragments of one or both HIV gp120 polypeptide sequences can be substituted for one or both of the corresponding HIV gp120 polypeptide sequences.

Preferably, the first gp120 polypeptide sequence contains neutralizing epitopes found in one or more gp120 polypeptides present in isolates from the geographical region where the initial vaccine (i.e., the vaccine that gives rise to the breakthrough isolate) is administered. More preferably, the first gp120 polypeptide sequence contains at least one of the more common neutralizing epitopes for the region, and most preferably the first gp120 polypeptide sequence contains at least one of the three most common neutralizing epitopes.

gp120 polypeptide sequences suitable for use as the first gp120 polypeptide sequence include gp120 MN, the Thai isolate A244 sequence (hereinafter "gp120 A244"), gp120 MN-GNE6 (Sequence ID No. 31; also known in the art as "gp120 GNE6"), and gp120 MN-GNE8 (Sequence ID No. 33; also known in the art as "gp120 GNE8"), and the like. gp120 MN, gp120 MN-GNE6, and gp120 MN-GNE8 are especially preferred for use as the first gp120 polypeptide sequence in initial vaccines for North America. gp120 A244 is especially preferred for use as the first gp120 polypeptide sequence in initial vaccines for Thailand.

In a variation of this embodiment, the vaccine includes two different (i.e., first and second) gp120 polypeptide sequences, or fragments thereof, in combination with a breakthrough isolate HIV gp120 polypeptide sequence. The latter can be from a vaccinee vaccinated with either or both of the first and second HIV gp120 polypeptide sequences.

Exemplary vaccines include those containing

combinations of gp120 MN, gp120 A244, gp120 MN-GNE6 (Sequence ID No. 31), and gp120 MN-GNE8 (Sequence ID No. 33). Combinations of gp120 MN and gp120 A244 or gp120 MN-GNE8 (Sequence ID No. 33) with a breakthrough isolate HIV gp120 polypeptide sequence are especially preferred.

In vaccines containing gp120 MN, the breakthrough isolate HIV gp120 polypeptide sequence can be an HIV gp120 polypeptide sequence selected from the group consisting of Sequence ID Nos. 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, and 28, and fragments thereof.

The term "subunit vaccine" is used herein, as in the art, to refer to a viral vaccine that does not contain virus, but rather contains one or more viral proteins or fragments of viral proteins. As used herein, the term "multivalent", means that the vaccine contains gp120 from at least two HIV isolates having different amino acid sequences.

The term "breakthrough isolate" or "breakthrough virus" is used herein, as in the art, to refer to a virus isolated from a vaccinee.

The terms "amino acid sequence", "polypeptide sequence", and "polypeptide" are used interchangeably herein as in the art, as are the terms "nucleic acid sequence", "nucleotide sequence", and "oligonucleotide".

Polypeptides from Breakthrough Isolates

The gp120 polypeptides of this invention correspond to the amino acid sequences of seven breakthrough isolates which are illustrated below in Table 1. A polypeptide of this invention includes an HIV gp120 amino acid sequence illustrated in Table 1 (Sequence ID Nos. 1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, and 27) and fragments thereof. The polypeptides of this invention can include fused

sequences from two or more HIV gp120 or gp160 amino acid sequences.

The polypeptide can also be joined to another viral protein, such as a flag epitope amino acid sequence. The term "flag epitope" is used herein, as in the art, to denote an amino acid sequence that includes an epitope recognized by a monoclonal antibody. Flag epitopes facilitate using single monoclonal antibody affinity purification of a plurality of different recombinant proteins, each having the flag epitope recognized by the monoclonal antibody. Numerous amino acid sequences can function as flag epitopes. The N-terminal sequences of Herpes Simplex Virus Type 1 (HSV-1) glycoprotein D (gD-1) is conveniently used as the flag epitope and its use is described in detail in the examples. The flag epitope is conveniently fused to the N terminus of the HIV gp120 polypeptide sequence. Alternatively, however, monoclonal antibodies that recognize neutralizing epitopes in the rgp120 sequences can be used to affinity purify the amino acid sequences, and a flag epitope can be omitted.

In addition, various signal sequences can be joined to a polypeptide of this invention. Although rgp120 is secreted to some extent in HIV cultures, the amount of the envelope glycoprotein released from (secreted by) the host cells varies widely from strain to strain. Various signal sequences can be introduced into the polypeptide by joining a nucleotide sequence encoding the signal sequence to the nucleotide sequence encoding the rgp120 to facilitate secretion of rgp120 from the cells. For example, Chiron HIV gp120 polypeptides include a signal sequence from tissue plasminogen activator (TPA) that provides good secretion of rgp120. Additional signal sequences are well known and include the N-terminal domain of murine

leukemia virus surface protein gp70 described by Kayman et al., *J. Virol.* 68:400-410 (1984).

Table 1 illustrates the nucleotide and deduced amino acid sequences for two clones of each the seven breakthrough isolates of this invention. The clones are: C6.1; C6.5; C8.3; C8.6; C15.2; C15.3; C7.2; C7.10; C11.5; C11.7; C10.5; C10.7; C17.1; and C17.3. These sequences are SEQ. ID. NOs. 1-28, the first sequence number for each clone being the nucleotide sequence and the second being the amino acid sequence. The amino acid sequence for MN and the nucleotide and deduced amino acid sequences for MN-GNE6 and MN-GNE8 are illustrated in the sequence listing hereinafter. In the listing for MN-GNE6, a stop codon appears at amino acid residue position 51. This stop codon can be replaced with a codon encoding the corresponding amino acid from MN or MN-GNE8 or another isolate.

TABLE 1

CLONE C6.1

| | |
|----|---|
| | GGG GTA CCT GTG TGG AAG GAA GCA ACC ACC ACT CTA 36 |
| 5 | Gly Val Pro Val Trp Lys Glu Ala Thr Thr Leu |
| | 1 5 10 |
| | TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GAG GTG 75 |
| | Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val |
| | 15 20 25 |
| 10 | CAT AAT GTT TGG GCC ACA CAT GCT TGT GTA CCC ACA GAC 114 |
| | His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp |
| | 30 35 |
| | CCA AAC CCA CAA GAA ATG GTA TTG GAA AAT GTG ACA GAA 153 |
| | Pro Asn Pro Gln Glu Met Val Leu Glu Asn Val Thr Glu |
| 15 | 40 45 50 |
| | GAT TTT AAC ATG TGG AAA AAT GAC ATG GTA GAA CAG ATG 192 |
| | Asp Phe Asn Met Trp Lys Asn Asp Met Val Glu Gln Met |
| | 55 60 |
| | CAT GAG GAT ATA ATC AGT TTA TGG GAT CAA AGC CTA AAA 231 |
| 20 | His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys |
| | 65 70 75 |
| | CCA TGT GTA AAA TTA ACC CCA CTC TGT ATT ACT TTA AAT 270 |
| | Pro Cys Val Lys Leu Thr Pro Leu Cys Ile Thr Leu Asn |
| | 80 85 90 |
| 25 | TGC ACC AAT TGG AAG AAG AAT GAT ACT AAA ACT AAT AGT 309 |
| | Cys Thr Asn Trp Lys Lys Asn Asp Thr Lys Thr Asn Ser |
| | 95 100 |
| | AGT AGT ACT ACA ACT AAT AAT AGT AGT GCT ACA GCT AAT 348 |
| 30 | Ser Ser Thr Thr Thr Asn Asn Ser Ser Ala Thr Ala Asn |
| | 105 110 115 |
| | AGT AGT AGT ACT ACA ACT AAT AGT AGT TGG GCA GAG ATA 387 |
| | Ser Ser Ser Thr Thr Thr Asn Ser Ser Trp Gly Glu Ile |
| | 120 125 |
| | AAG GAG GGA GAA ATA AAG AAC TGC TCT TTC AAT ATC ACC 426 |
| 35 | Lys Glu Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile Thr |
| | 130 135 140 |
| | ACA AGC ATA AGA GAC AAG GTG AAG AAA GAA TAT CCA CTT 465 |
| | Thr Ser Ile Arg Asp Lys Val Lys Lys Glu Tyr Ala Leu |
| | 145 150 155 |
| 40 | TTT TAT AGC CTT GAT GTA GTA CCA ATA GAA AAT GAT AAT 504 |
| | Phe Tyr Ser Leu Asp Val Val Pro Ile Glu Asn Asp Asn |
| | 160 165 |
| | ACT AGC TAT AGG TTG AGA AGT TGT AAC ACC TCA GTC ATT 543 |
| | Thr Ser Tyr Arg Leu Arg Ser Cys Asn Thr Ser Val Ile |
| 45 | 170 175 180 |
| | ACA CAA GCC TGT CCA AAG GTA ACT TTT GAG CCA ATT CCC 582 |
| | Thr Gln Ala Cys Pro Lys Val Thr Phe Glu Pro Ile Pro |
| | 185 190 |
| | ATA CAT TAT TGT ACC CCG GCT GGT TTT GCG ATT CTG AAG 621 |
| 50 | Ile His Tyr Cys Thr Pro Ala Gly Phe Ala Ile Leu Lys |
| | 195 200 205 |
| | TGT AGA GAT AAA AAG TTC AAT GGA ACA GGA CCA TGC AAA 660 |
| | Cys Arg Asp Lys Lys Phe Asn Gly Thr Gly Pro Cys Lys |
| | 210 215 220 |
| 55 | AAT GTT AGC ACA GTA CAA TGT GCA CAT GGA ATT AAG CCA 699 |
| | Asn Val Ser Thr Val Gln Cys Ala His Gly Ile Lys Pro |
| | 225 230 |
| | GTA GTG TCA ACT CAA CTG CTG TTA AAT GGC AGC CTA GCA 738 |
| | Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala |
| 60 | 235 240 245 |
| | GAA GAA GAG GTA ATA ATT AGA TCT GCC AAT TTC TCA AAC 777 |
| | Glu Glu Glu Val Ile Ile Arg Ser Ala Asn Phe Ser Asn |
| | 250 255 |

AAT GCT AAA ATC ATA ATA GTA CAG TTG AGG GAA CCT GTA 816
 Asn Ala Lys Ile Ile Ile Val Gln Leu Arg Glu Pro Val
 260 265 270
 5 GAA ATT AAT TGT ACA AGA CCC AGC AAC AAT ACA ATA AAA 855
 Glu Ile Asn Cys Thr Arg Pro Ser Asn Asn Thr Ile Lys
 275 280 285
 GGT ATA CAC ATA GGA CCA GGG AGA GCA TTT TAT GCA ACA 894
 Gly Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala Thr
 290 295
 10 GGA GAC ATA CGA GGA GAT ATA AGA CAA GCA CAT TGT AAC 933
 Gly Asp Ile Arg Gly Asp Ile Arg Gln Ala His Cys Asn
 300 305 310
 ATT AGT GGA GCA AAA TGG AAT AAC ACT TTA AAG AAG GTA 972
 Ile Ser Gly Ala Lys Trp Asn Asn Thr Leu Lys Lys Val
 315 320
 15 GTT AAA AAA TTA AAA GAA CAA TTT CCA AAT AAA ACA ATA 1011
 Val Lys Lys Leu Lys Glu Gln Phe Pro Asn Lys Thr Ile
 325 330 335
 GTC TTT AAC CAT TCC TCA GGA GGG GAC CCA GAA ATT GTA 1050
 Val Phe Asn His Ser Ser Gly Gly Asp Pro Glu Ile Val
 340 345 350
 20 ATG CAC AGT TTT AAT TGT CAA GGG GAA TTT TTC TAC TGT 1089
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 365 370 375
 ACA GAG TCA AAT AAC AAT GAT AGT ACT ATT ACA CTC CCA 1167
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 380 385
 30 TGC AGA ATA AAA CAA ATT ATA AAC ATG TGG CAG GAA ATA 1206
 Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Ile
 390 395 400
 GGA AAA GCA ATG TAT GCC CCT CCC ACC AGA GGA GAA ATT 1245
 Gly Lys Ala Met Tyr Ala Pro Pro Thr Arg Gly Glu Ile
 405 410 415
 35 AAA TGT TCA TCA AAT ATT ACA GGA CTA CTG TTA ATA AGA 1284
 Lys Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Ile Arg
 420 425
 40 GAT GGT GGT ATT AAC ACT AGC GAT GCC ACC GAG ACC TTC 1323
 Asp Gly Gly Ile Asn Thr Ser Asp Ala Thr Glu Thr Phe
 430 435 440
 AGA CCG GGA GGA GCA GAT ATG AGG GAC AAT TGG AGA AGT 1362
 Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser
 445 450
 45 GAA TTA TAT AAA TAT AAA GTA GTG AAA ATT GAG CCA TTA 1401
 Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu
 455 460 465
 GGA GTA GCA CCC ACC AAG GCA AAG AGA AGA GTG GTG CAG 1440
 Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 470 475 480
 50 AGA GAA AAA AGA GCA GTA ACA CTA GGA GCT ATG TTC CTT 1479
 Arg Glu Lys Arg Ala Val Thr Leu Gly Ala Met Phe Leu
 485 490
 55 GGG TTC TTA GGA GCA TAA AGC TTC 1503
 Gly Phe Leu Gly Ala Xaa Ser Phe
 495 500 501

CLONE C6.5

60 GGG GTA CCT GTA TGG AAA GAA GCA ACC ACC ACT CTA 36
 Gly Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu
 1 5 10
 TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GAG GTG 75
 Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val
 15 20 25
 65

| | | |
|----|---|-----|
| | CAT AAT GTT TGG GCC ACA CAT GCT TGT GTA CCC AGA GAC | 114 |
| | His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp | |
| | 30 35 | |
| 5 | CCA AAC CCA CAA GAA ATG GTA TTG GAA AAT GTG ACA GAA | 153 |
| | Pro Asn Pro Gln Glu Met Val Leu Glu Asn Val Thr Glu | |
| | 40 45 50 | |
| | GAT TTT AAC ATG TGG AAA AAT GAC ATG GTA GAA CAG ATG | 192 |
| | Asp Phe Asn Met Trp Lys Asn Asp Met Val Glu Gln Met | |
| | 55 60 | |
| 10 | CAT GAG ANT ATA ATC AGT TTA TGG GAT CAA AGC CTA AAA | 231 |
| | His Glu Xaa Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys | |
| | 65 70 75 | |
| | CCA TGT GTA AAA TTA ACC CCA CTC TGT ATT ACT TTA AAT | 270 |
| | Pro Cys Val Lys Leu Thr Pro Leu Cys Ile Thr Leu Asn | |
| 15 | 80 85 90 | |
| | TGC ACC AAT TGG AAG GAG AAT GAT ACT AAA ACT AAT AGT | 309 |
| | Cys Thr Asn Trp Lys Glu Asn Asp Thr Lys Thr Asn Ser | |
| | 95 100 | |
| 20 | AGT AGT ACT ACA ACT AAT AAT AGT AGT GCT ACA GCT AAT | 348 |
| | Ser Ser Thr Thr Thr Asn Asn Ser Ser Ala Thr Ala Asn | |
| | 105 110 115 | |
| | AGT AGT AGT ACT ACA ACT AAT AGT AGT TGG GGA GAG ATA | 387 |
| | Ser Ser Ser Thr Thr Thr Asn Ser Ser Trp Gly Glu Ile | |
| | 120 125 | |
| 25 | AAG GAG GGA GAA ATA AAG AAC TGC TCT TTC AAT ATC ACC | 426 |
| | Lys Glu Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile Thr | |
| | 130 135 140 | |
| | ACA GGC ATA AGA GAC AAG GTG AAG AAA GAA TAT GCA CTT | 465 |
| | Thr Gly Ile Arg Asp Lys Val Lys Lys Glu Tyr Ala Leu | |
| 30 | 145 150 155 | |
| | TTT TAT AGC CTT GAT GTA GTA CCA ATA GAA AAT GAT AAT | 504 |
| | Phe Tyr Ser Leu Asp Val Val Pro Ile Glu Asn Asp Asn | |
| | 160 165 | |
| 35 | ACT AGC TAT AGG TTG AGA AGT TGT AAC ACC TCA GTC ATT | 543 |
| | Thr Ser Tyr Arg Leu Arg Ser Cys Asn Thr Ser Val Ile | |
| | 170 175 180 | |
| | ACA CAA GCC TGT CCA AAG GTA ACT TTT GAG CCA ATT CCC | 582 |
| | Thr Gln Ala Cys Pro Lys Val Thr Phe Glu Pro Ile Pro | |
| | 185 190 | |
| 40 | ATA CAT TAT TGT ACC CCG GCT GGT TTT GCG ATT CTG AAG | 621 |
| | Ile His Tyr Cys Thr Pro Ala Gly Phe Ala Ile Leu Lys | |
| | 195 200 205 | |
| | TGT AAA GAT AAA AAG TTC AAT GGA ACA GGA CCA TGC AAA | 660 |
| | Cys Lys Asp Lys Lys Phe Asn Gly Thr Gly Pro Cys Lys | |
| 45 | 210 215 220 | |
| | AAT GTT AGC ACA GTA CAA TGT ACA CAT GGA ATT AAG CCA | 699 |
| | Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Lys Pro | |
| | 225 230 | |
| 50 | GTA GTG TCA ACT CAA CTG CTG TTA AAT GGC AGC CTA GCA | 738 |
| | Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala | |
| | 235 240 245 | |
| | GAA GAA GAG GTA ATA ATT AGA TCT GCC AAT TTC TCA AAC | 777 |
| | Glu Glu Glu Val Ile Ile Arg Ser Ala Asn Phe Ser Asn | |
| | 250 255 | |
| 55 | AAT GCT AAA ATC ATA ATA GTA CAG TTG AAG GAA CCT GTA | 816 |
| | Asn Ala Lys Ile Ile Ile Val Gln Leu Lys Glu Pro Val | |
| | 260 265 27 | |
| | GAA ATT AAT TGT ACA AGA CCC AGC AAC AAT ACA ATA AAA | 855 |
| | Glu Ile Asn Cys Thr Arg Pro Ser Asn Asn Thr Ile Lys | |
| 60 | 275 280 285 | |
| | GGT ATA CAC ATA GGA CCA GGG AGA GCA TTT TAT GCA ACA | 894 |
| | Gly Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala Thr | |
| | 290 295 | |

GGA GAC ATA CGA GGA GAT ATA AGA CAA GCA CAT TGT AAC 933
 Gly Asp Ile Arg Gly Asp Ile Arg Gln Ala His Cys Asn
 300 305 310
 ATT AGT GGA GCA AAA TGG AAT AAC ACT TTA AAG AAG GTA 972
 Ile Ser Gly Ala Lys Trp Asn Asn Thr Leu Lys Lys Val
 315 320
 GTT ATA AAA TTA AAA GAA CAA TTT CCA AAT AAA ACA ATA 1011
 Val Ile Lys Leu Lys Glu Gln Phe Pro Asn Lys Thr Ile
 325 330 335
 10 GTC TTT AAC CAT TCC TCA GGA GGG GAC CCA GAA ATT GTA 1050
 Val Phe Asn His Ser Ser Gly Gly Asp Pro Glu Ile Val
 340 345 350
 ATG CAC AGT TTT AAT TGT CAA GGG GAA TTT TTC TAC TGT 1089
 Met His Ser Phe Asn Cys Gln Gly Glu Phe Phe Tyr Cys
 355 360
 15 AAT ACA ACG AAG CTG TTT AAT AGT ACT TGG AAT GAT ACT 1128
 Asn Thr Thr Lys Leu Phe Asn Ser Thr Trp Asn Asp Thr
 365 370 375
 ACA GAG TCA AAT AAC AAT GAT AGT ACT ATT ACA CTC CCA 1167
 Thr Glu Ser Asn Asn Asp Ser Thr Ile Thr Leu Pro
 380 385
 TGC AGA ATA AAA CAA ATT ATA AAC ATG TGG CAG GAA GTA 1206
 Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val
 390 395 400
 25 GGA AAA GCA ATG TAT GCC CCT CCC ATC AGA GGA GAA ATT 1245
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Glu Ile
 405 410 415
 AAA TGT TCA TCA AAT ATT ACA GGA CTA CTG TTA ACA AGA 1284
 Lys Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg
 420 425
 30 GAT GGT GGT ATT AAC ACT AGC GAT GCC ACC GAG ACC TTC 1323
 Asp Gly Gly Ile Asn Thr Ser Asp Ala Thr Glu Thr Phe
 430 435 440
 AGA CCG GGA GGA GGA GAT ATG AGG GAC AAT TGG AGA ACT 1362
 Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser
 445 450
 GAA TTA TAT AAA TAT AAA GTA GTG AAA ATT GAG CCA TTA 1401
 Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu
 455 460 465
 40 GGA GTA GCA CCC ACC AAG GCA AAG AGA AGA GTG GTG CAG 1440
 Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 470 475 480
 AGA GAA AAA AGA GCA GTA ACA CTA GGA GCT ATG TTC CTT 1479
 Arg Glu Lys Arg Ala Val Thr Leu Gly Ala Met Phe Leu
 485 490
 45 GGG TTC TTG GGA GCA TAA AGC TTC 1503
 Gly Phe Leu Gly Ala Xaa Ser Phe
 495 500 501
 50 CLONE C8.3
 C GTA CCT GTA TGG AAA GAA GCA ACC ACC ACT CTA TTT 37
 Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe
 1 5 10
 TGT GCA TCA GAT GCT AAA GCA TAT GAT ACA GAG GTA CAT 76
 Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val His
 15 20 25
 AAT GTT TGG GCT ACA CAT GCC TGT GTA CCC ACA GAC CCC 115
 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro
 30 35
 60 AAC CCA CAA GAA GTA GTA TTG GAA AAT GTA ACA GAA AAT 154
 Asn Pro Gln Glu Val Val Leu Glu Asn Val Thr Glu Asn
 40 45 50
 TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAG ATG CAT 193
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His
 55 60

| | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|------|
| | GAG | GAT | ATA | ATC | AGT | TTA | TGG | GAT | CAA | AGT | CTA | AAG | CCA | 232 |
| | Glu | Asp | Ile | Ile | Ser | Leu | Trp | Asp | Gln | Ser | Leu | Lys | Pro | |
| | 65 | | | | | 70 | | | | | 75 | | | |
| 5 | TGT | GTA | AAA | TTA | ACC | CCA | CTC | TGT | GTT | ACT | TTA | AAT | TGC | 271 |
| | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Val | Thr | Leu | Asn | Cys | |
| | | | 80 | | | | | 85 | | | | | 90 | |
| | ACT | AAT | TTG | GAG | AAT | GCT | AAT | AAT | ACC | GAG | AAT | GCT | AAT | 310 |
| | Thr | Asn | Leu | Glu | Asn | Ala | Asn | Asn | Thr | Glu | Asn | Ala | Asn | |
| | | | | | 95 | | | | | 100 | | | | |
| 10 | AAT | ACC | AAT | AAT | TAT | ACC | TTG | GGG | ATG | GAG | AGA | GGT | GAA | 349 |
| | Asn | Thr | Asn | Asn | Tyr | Thr | Leu | Gly | Met | Glu | Arg | Gly | Glu | |
| | | | | | | 110 | | | | | | 115 | | |
| | ATA | AAA | AAC | TGC | TCT | TTC | AAT | ATC | ACC | ACA | AGC | TTA | AGA | 388 |
| | Ile | Lys | Asn | Cys | Ser | Phe | Asn | Ile | Thr | Thr | Ser | Leu | Arg | |
| 15 | | | | | 120 | | | | | | | | | |
| | GAT | AAG | GTG | AAA | AAA | GAA | TAT | GCA | TTG | TTT | TAT | AAA | CTT | 427 |
| | Asp | Lys | Val | Lys | Lys | Glu | Tyr | Ala | Leu | Phe | Tyr | Lys | Leu | |
| | | | | | | 135 | | | | | | 140 | | |
| 20 | GAT | GTA | GTA | CAA | ATA | GAT | AAT | AGT | ACC | AAC | TAT | AGG | CTG | 466 |
| | Asp | Val | Val | Gln | Ile | Asp | Asn | Ser | Thr | Asn | Tyr | Arg | Leu | |
| | | | | | | 145 | | | | | | | 155 | |
| | ATA | AGT | TGT | AAT | ACC | TCA | GTC | ATT | ACA | CAG | GCC | TGT | CCA | 505 |
| | Ile | Ser | Cys | Asn | Thr | Ser | Val | Ile | Thr | Gln | Ala | Cys | Pro | |
| | | | | | 160 | | | | | | 165 | | | |
| 25 | AAG | GTA | TCC | TTT | GAG | CTA | ATT | CCC | ATA | CAT | TAT | TGT | GCC | 544 |
| | Lys | Val | Ser | Phe | Glu | Leu | Ile | Pro | Ile | His | Tyr | Cys | Ala | |
| | | | | | | 170 | | | | | | 180 | | |
| | CCG | GCT | GGT | TTT | GCG | ATT | CTA | AAG | TGT | AAA | GAT | AAG | AAG | 583 |
| | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Lys | Asp | Lys | Lys | |
| 30 | | | | | 185 | | | | | | | | | |
| | TTC | AAT | GGA | ACA | GGA | CCA | TGT | AAA | AAT | GTC | AGC | ACA | GTA | 622 |
| | Phe | Asn | Gly | Thr | Gly | Pro | Cys | Lys | Asn | Val | Ser | Thr | Val | |
| | | | | | | 195 | | | | | | 205 | | |
| 35 | CAA | TGT | ACA | CAT | GGA | ATT | AGA | CCA | GTA | GTA | TCA | ACT | CAA | 661 |
| | Gln | Cys | Thr | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | Gln | |
| | | | | | 210 | | | | | | | | 220 | |
| | CTA | CTG | TTA | AAT | GGC | AGT | CTA | GCA | GAA | GAA | GAG | ATA | GTA | 700 |
| | Leu | Leu | Leu | Asn | Gly | Ser | Leu | Ala | Glu | Glu | Glu | Ile | Val | |
| | | | | | 225 | | | | | | 230 | | | |
| 40 | ATT | AGA | TCT | GAA | AAT | ATC | ACA | GAC | AAT | GCT | AAA | ACC | ATA | 739 |
| | Ile | Arg | Ser | Glu | Asn | Ile | Thr | Asp | Asn | Ala | Lys | Thr | Ile | |
| | | | | | | 235 | | | | | | 245 | | |
| | ATA | GTG | CAG | CTA | AAT | GAA | TCT | ATA | GTG | ATT | AAT | TGT | ACA | 778 |
| | Ile | Val | Gln | Leu | Asn | Glu | Ser | Ile | Val | Ile | Asn | Cys | Thr | |
| 45 | | | | | 250 | | | | | | | | | |
| | AGA | CCC | AAT | AAC | AAC | ACA | AGA | AAA | AGT | ATA | AAT | ATA | GGA | 817 |
| | Arg | Pro | Asn | Asn | Asn | Thr | Arg | Lys | Ser | Ile | Asn | Ile | Gly | |
| | | | | | | 260 | | | | | | 270 | | |
| 50 | CCA | GGG | AGA | GCA | TTC | TAT | ACA | ACA | GGA | GAC | ATA | ATA | GGA | 856 |
| | Pro | Gly | Arg | Ala | Phe | Tyr | Thr | Thr | Gly | Asp | Ile | Ile | Gly | |
| | | | | | | 275 | | | | | | | 285 | |
| | GAT | ATA | AGA | CAA | GCA | CAT | TGT | AAC | CTT | AGT | AAA | ACA | CAA | 895 |
| | Asp | Ile | Arg | Gln | Ala | His | Cys | Asn | Leu | Ser | Lys | Thr | Gln | |
| | | | | | 290 | | | | | | 295 | | | |
| 55 | TGG | GAA | AAA | ACG | TTA | AGA | CAG | ATA | GCT | ATA | AAA | TTA | GAA | 934 |
| | Trp | Glu | Lys | Thr | Leu | Arg | Gln | Ile | Ala | Ile | Lys | Leu | Glu | |
| | | | | | | 300 | | | | | | 310 | | |
| | GAA | AAA | TTT | AAG | AAT | AAA | ACA | ATA | GCC | TTT | AAT | AAA | TCC | 973 |
| | Glu | Lys | Phe | Lys | Asn | Lys | Thr | Ile | Ala | Phe | Asn | Lys | Ser | |
| 60 | | | | | 315 | | | | | | | | | |
| | TCA | GGA | GGG | GAC | CCA | GAA | ATT | GTA | ATG | CAC | AGT | TTT | AAT | 1012 |
| | Ser | Gly | Gly | Asp | Pro | Glu | Ile | Val | Met | His | Ser | Phe | Asn | |
| | | | | | 325 | | | | | | | 335 | | |

TGT GGA GGG GAA TTT TTC TAC TGT AAT ACA ACA AAA CTG 1051
 Cys Gly Gly Glu Ph Phe Tyr Cys Asn Thr Thr Lys Leu
 340 345 350
 5 TTT AAT AGT ACC TGG AAT TTA ACA CAA CCG TTT AGT AAT 1090
 Phe Asn Ser Thr Trp Asn Leu Thr Gln Pro Phe Ser Asn
 355 360
 ACC GGG AAT CGT ACT GAA GAG TTA AAT ATT ACA CTC CCA 1129
 Thr Gly Asn Arg Thr Glu Glu Leu Asn Ile Thr Leu Pro
 365 370 375
 10 TGC AGA ATA AAA CAA ATC ATA AAC TTG TGG CAG GAA GTA 1168
 Cys Arg Ile Lys Gln Ile Ile Asn Leu Trp Gln Glu Val
 380 385
 GGC AAA GCA ATG TAT GCC CCT CCC ATC AGA GGA CAA ATT 1207
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile
 390 395 400
 15 AGA TGT TCA TCA AAT ATT ACA GGG CTA CTA TTA ACA AGA 1246
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg
 405 410 415
 20 GAT GGT GGA AGT AAC ACC GGT GAC AAC AGG ACT GAG ACC 1285
 Asp Gly Gly Ser Asn Thr Gly Asp Asn Arg Thr Glu Thr
 420 425
 TTT AGA CCT GGA GGA GGA GAT ATG AGG GAC AAT TGG AGA 1324
 Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 430 435 440
 25 AGT GAA TTA TAT AAA TAT AAA GTA GTA AGA ATT GAA CCA 1363
 Ser Glu Leu Tyr Lys Tyr Lys Val Val Arg Ile Glu Pro
 445 450
 TTA GGA GTA GCA CCC ACC CAG GCA AAG AGA AGA GTG GTG 1402
 Leu Gly Val Ala Pro Thr Gln Ala Lys Arg Arg Val Val
 455 460 465
 30 CAA AGA GAA AAA AGA GCA GTG GGG ATA GGA GCT ATG TTC 1441
 Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Met Phe
 470 475 480
 CTT GGG TTC TTG GGA GAT AA 1461
 35 Leu Gly Phe Leu Gly Asp
 485 486

CLONE C8.6

G GTA CCT GTG TGG AAA GAA GCA ACC ACC ACT CTA TTT 37
 40 Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe
 1 5 10
 TGT GCA TCA GAT GCT AAA GCA TAT GAT ACA GAG GTA CAT 76
 Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val His
 15 20 25
 45 AAT GTT TGG GCT ACA CAT GCC TGT GTA CCC ACA GAC CCC 115
 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro
 30 35
 AAC CCA CAA GAA GTA GTA TTG GAA AAT GTA ACA GAA AAT 154
 Asn Pro Gln Glu Val Val Leu Glu Asn Val Thr Glu Asn
 40 45 50
 50 TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAG ATG CAT 193
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His
 55 60
 GAG GAT ATA ATC AGT TTA TGG GAT CAA AGT CTA AAG CCA 232
 Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys Pro
 65 70 75
 55 TGT GTA AAA TTA ACC CCA CTC TGT GTT ACT TTA AAT TGC 271
 Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys
 80 85 90
 60 ACT AAT TTG GAG AAT GCT AAT AAT ACC GAG AAT GCT AAT 310
 Thr Asn Leu Glu Asn Ala Asn Asn Thr Glu Asn Ala Asn
 95 100
 AAT ACC AAT AAT TAT ACC TTG GGG ATG GAG AGA GGT GAA 349
 Asn Thr Asn Asn Tyr Thr Leu Gly Met Glu Arg Gly Glu
 105 110 115
 65

| | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|------|
| | AGA | AAA | AAC | TGC | TCT | TTC | AAT | ATC | ACC | ACA | AGC | TTA | AGA | 388 |
| | Arg | Lys | Asn | Cys | Ser | Phe | Asn | Ile | Thr | Thr | Ser | Leu | Arg | |
| | | | | 120 | | | | | 125 | | | | | |
| 5 | GAT | AAG | GGG | AAA | AAA | GAA | TAT | GCA | TTG | TTT | TAT | AAA | CTT | 427 |
| | Asp | Lys | Gly | Lys | Lys | Glu | Tyr | Ala | Leu | Phe | Tyr | Lys | Leu | |
| | | | | 130 | | | | | 135 | | | | 140 | |
| | GAT | GTA | GTA | CAA | ATA | GAT | AAT | AGT | ACC | AAC | TAT | AGG | CTG | 466 |
| | Asp | Val | Val | Gln | Ile | Asp | Asn | Ser | Thr | Asn | Tyr | Arg | Leu | |
| | | | | 145 | | | | | 150 | | | | 155 | |
| 10 | ATA | AGT | TGT | AAT | ACC | TCA | GTC | ATT | ACA | CAG | GCC | TGT | CCA | 505 |
| | Ile | Ser | Cys | Asn | Thr | Ser | Val | Ile | Thr | Gln | Ala | Cys | Pro | |
| | | | | | 160 | | | | | | | | 165 | |
| | AAG | GTA | TCC | TTT | GAG | CCA | ATT | CCC | ATA | CAT | TAT | TGT | GCC | 544 |
| | Lys | Val | Ser | Phe | Glu | Pro | Ile | Pro | Ile | His | Tyr | Cys | Ala | |
| 15 | | | | 170 | | | | | 175 | | | | 180 | |
| | CCG | GCT | GGT | TTT | GCG | ATT | CTA | AAG | TGT | AAA | GAT | AAG | AAG | 583 |
| | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Lys | Asp | Lys | Lys | |
| | | | | 185 | | | | | 190 | | | | | |
| 20 | TTC | AAT | GGA | ACA | GGA | CCA | TGT | AAA | AAT | GTC | AGG | ACA | GTA | 622 |
| | Phe | Asn | Gly | Thr | Gly | Pro | Cys | Lys | Asn | Val | Arg | Thr | Val | |
| | | | | 195 | | | | | 200 | | | | 205 | |
| | CAA | TGT | ACA | CAT | GGA | ATT | AGA | CCA | GTA | GTA | TCA | ACT | CAA | 661 |
| | Gln | Cys | Thr | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | Gln | |
| | | | | 210 | | | | | 215 | | | | 220 | |
| 25 | CTA | CTG | TTA | AAT | GGC | AGT | CTA | GCA | GAA | GAA | GAG | ATA | GTA | 700 |
| | Leu | Leu | Leu | Asn | Gly | Ser | Leu | Ala | Glu | Glu | Glu | Ile | Val | |
| | | | | | 225 | | | | | | 230 | | | |
| | ATT | AGA | TCT | GAA | AAT | ATC | ACA | GAC | AAT | GCT | AAA | ACC | ATA | 739 |
| | Ile | Arg | Ser | Glu | Asn | Ile | Thr | Asp | Asn | Ala | Lys | Thr | Ile | |
| 30 | | | | 235 | | | | | 240 | | | | 245 | |
| | ATA | GTG | CAG | CTA | AAT | GAA | TCT | ATA | GTG | ATT | AAT | TGT | ACA | 778 |
| | Ile | Val | Gln | Leu | Asn | Glu | Ser | Ile | Val | Ile | Asn | Cys | Thr | |
| | | | | 250 | | | | | 255 | | | | | |
| 35 | AGA | CCC | AAT | AAC | AAC | ACA | AGA | AAA | AGT | ATA | AAT | ATA | GGA | 817 |
| | Arg | Pro | Asn | Asn | Asn | Thr | Arg | Lys | Ser | Ile | Asn | Ile | Gly | |
| | | | | 260 | | | | | 265 | | | | 270 | |
| | CCA | GGG | AGA | GCA | TTC | TAT | ACA | ACA | GGA | GAC | ATA | ATA | GGA | 856 |
| | Pro | Gly | Arg | Ala | Phe | Tyr | Thr | Thr | Gly | Asp | Ile | Ile | Gly | |
| | | | | 275 | | | | | 280 | | | | 285 | |
| 40 | GAT | ATA | AGA | CAA | GCA | CAT | TGT | AAC | CTT | AGT | AAA | ACA | CAA | 895 |
| | Asp | Ile | Arg | Gln | Ala | His | Cys | Asn | Leu | Ser | Lys | Thr | Gln | |
| | | | | 290 | | | | | 295 | | | | | |
| | TGG | GAA | AAA | ACG | TTA | AGA | CAG | ATA | GCT | ATA | AAA | TTA | GAA | 934 |
| | Trp | Glu | Lys | Thr | Leu | Arg | Gln | Ile | Ala | Ile | Lys | Leu | Glu | |
| 45 | | | | 300 | | | | | 305 | | | | 310 | |
| | GAA | AAA | TTT | AAG | AAT | AAA | ACA | ATA | GCC | TTT | AAT | AAA | TCC | 973 |
| | Glu | Lys | Phe | Lys | Asn | Lys | Thr | Ile | Ala | Phe | Asn | Lys | Ser | |
| | | | | 315 | | | | | 320 | | | | | |
| 50 | TCA | GGA | GGG | GAC | CCA | GAA | ATT | GTA | ATG | CAC | AGT | TTT | AAT | 1012 |
| | Ser | Gly | Gly | Asp | Pro | Glu | Ile | Val | Met | His | Ser | Phe | Asn | |
| | | | | 325 | | | | | 330 | | | | 335 | |
| | TGT | GGA | GGG | GGA | TTT | TTC | TAC | TGT | AGT | ACG | AGA | AAA | CTG | 1051 |
| | Cys | Gly | Gly | Gly | Phe | Phe | Tyr | Cys | Ser | Thr | Arg | Lys | Leu | |
| | | | | 340 | | | | | 345 | | | | 350 | |
| 55 | TTT | AAT | AGT | ACC | TGG | AAT | TTA | ACA | CAA | CCG | TTT | AGT | AAT | 1090 |
| | Phe | Asn | Ser | Thr | Trp | Asn | Leu | Thr | Gln | Pro | Phe | Ser | Asn | |
| | | | | | 355 | | | | 360 | | | | | |
| | ACC | GGG | GAT | CGT | ACT | GAA | GAG | TTA | AAT | ATT | ACA | CTC | CCA | 1129 |
| | Thr | Gly | Asp | Arg | Thr | Glu | Glu | Leu | Asn | Ile | Thr | Leu | Pro | |
| 60 | | | | 365 | | | | | 370 | | | | 375 | |
| | TGC | AGA | ATA | AAA | CAA | ATC | ATA | AAC | TTG | TGG | CAG | GAA | GTA | 1168 |
| | Cys | Arg | Ile | Lys | Gln | Ile | Ile | Asn | Leu | Trp | Gln | Glu | Val | |
| | | | | 380 | | | | | 385 | | | | | |

GGC AAA GCA ATG TAT GCC CCT CCC ATC AGA GGA CAA ATT 1207
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile
 390 395 400
 5 AGA TGT TCA TCA AAT ATT ACA GGG CTA CTA TTA AGG AGA 1246
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Arg Arg
 405 410 415
 GAT GGT GGA AGT AAC ACC AGT GAC AAC CAG ACT GAG ACC 1285
 Asp Gly Gly Ser Asn Thr Ser Asp Asn Gln Thr Glu Thr
 420 425
 10 TTT AGA CCT GGG GGA GGA GAT ATG AGG GAC AAG TGG ACA 1324
 Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Lys Trp Arg
 430 435 440
 AGT GAA TTA TAT AAA TAT AAA GTA GTA AGA ATT GAA CCA 1363
 Ser Glu Leu Tyr Lys Tyr Lys Val Val Arg Ile Glu Pro
 445 450
 15 TTA GGA GTA GCA CCC ACC CAG GCA AAG AGA AGA GTG GTG 1402
 Leu Gly Val Ala Pro Thr Gln Ala Lys Arg Arg Val Val
 455 460 465
 CAA AGA GAA AAA AGA GCA GTG GGG ATA GGA GCT ATG TTC 1441
 Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Met Phe
 470 475 480
 CTT AGG TTC TTA GGA GAT AAA GCT TCT AGA GTC 1474
 Leu Arg Phe Leu Gly Asp Lys Ala Ser Arg Val
 485 490 491
 25
CLONE C15.2
 CTC GAG GTA CCT GTA TGG AAA GAA GCA ACT ACC ACT 36
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr
 1 5 10
 30 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT AAT ACA GAG 75
 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asn Thr Glu
 15 20 25
 AAA CAT AAT GTT TGG GCC ACA CAC GCC TGT GTA CCC ACA 114
 Lys His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr
 30 35
 35 GAT CCC AAC CCA CAA GAA GTA GTA TTG GGA AAT GTG ACA 153
 Asp Pro Asn Pro Gln Glu Val Val Leu Gly Asn Val Thr
 40 45 50
 GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA 192
 Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln
 55 60
 ATG CAT GAA GAT ATA ATC AGT TTA TGG GAT CAA AGT CTA 231
 Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu
 65 70 75
 45 AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT GTT ACT TTA 270
 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu
 80 85 90
 AAT TGC ACT GAT GAT TTA GGG AAT GCT ACT AAT ACC AAT 309
 Asn Cys Thr Asp Asp Leu Gly Asn Ala Thr Asn Thr Asn
 95 100
 50 AGT AGT GCC ACT ACC AAT AGT AGT AGT TGG GAA GAA ATG 348
 Ser Ser Ala Thr Thr Asn Ser Ser Ser Trp Glu Glu Met
 105 110 115
 AAG GGG GAA ATG AAA AGA TGC TCT TTC AAT ATC ACC ACA 387
 Lys Gly Glu Met Lys Arg Cys Ser Phe Asn Ile Thr Thr
 120 125
 AGC ATA AGA GAT AAG ATT AAG AAA GAA CAT GCA CTT TTC 426
 Ser Ile Arg Asp Lys Ile Lys Lys Glu His Ala Leu Phe
 130 135 140
 60 TAT AGA CTT GAT GTA CCA ATA GAT AAT GAT AAT ACC 465
 Tyr Arg Leu Asp Val Val Pro Ile Asp Asn Asp Asn Thr
 145 150 155

ACA TAT AGG TTG ATA AAT TGT AAT ACC TCA GTC ATT ACA 504
 Thr Tyr Arg Leu Ile Asn Cys Asn Thr Ser Val Ile Thr
 160 165
 5 CAG GCC TGT CCA AAG GTA TCA TTT GAG CCA ATT CCC ATA 543
 Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile
 170 175 180
 CAT TTT TGT GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT 582
 His Phe Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys
 185 190
 10 AAT AAT AAG ACG TTC GAG GGA AAA GGA CCA TGT AAA AAT 621
 Asn Asn Lys Thr Phe Glu Gly Lys Gly Pro Cys Lys Asn
 195 200 205
 GTC AGT ACA GTA CAA TGC ACA CAT GGA ATT AGG CCA GTA 660
 Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val
 210 215 220
 15 GTG TCA ACT CAA CTG CTG TTA AAT GGC AGT CTA GCA GAA 699
 Val Ser Thr Gln Leu Leu Asn Gly Ser Leu Ala Glu
 225 230
 20 GAA GAG GTA ATA ATT AGA TCT GAC AAT ATC ACA GAC AAT 738
 Glu Glu Val Ile Ile Arg Ser Asp Asn Ile Thr Asp Asn
 235 240 245
 ACT AAA ACC ATT ATA GTA CAG CTA AAC GAA TCT GTA GTA 777
 Thr Lys Thr Ile Ile Val Gln Leu Asn Glu Ser Val Val
 250 255
 25 ATT AAT TGT ACA AGA CCC AAC AAC AAT ACA AGA AAA AGT 816
 Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg Lys Ser
 260 265 270
 30 ATA CAT ATA GGA CCA GGG AGT GCA TTT TTT GCA ACA GGA 855
 Ile His Ile Gly Pro Gly Ser Ala Phe Phe Ala Thr Gly
 275 280 285
 GAA ATA ATA GGA GAT ATA AGA CAA GCA CAC TGT AAC CTT 894
 Glu Ile Ile Gly Asp Ile Arg Gln Ala His Cys Asn Leu
 290 295
 35 AGT AGA ACA CAA TGG AAT AAC ACT TTA GGA AAG ATA GTC 933
 Ser Arg Thr Gln Trp Asn Asn Thr Leu Gly Lys Ile Val
 300 305 310
 ATA AAA TTA AGA GAA CAA TTT AGA AAA CAA TTT GGA GAA 972
 Ile Lys Leu Arg Glu Gln Phe Arg Lys Gln Phe Gly Glu
 315 320
 40 AAA ACA ATA GTC TTT AAT CGA TCC TCA GGA GGG GAC CCG 1011
 Lys Thr Ile Val Phe Asn Arg Ser Ser Gly Gly Asp Pro
 325 330 335
 GAA ATT GCA ATG CAC AGT TTT AAT TGT GGA GGG GAA TTT 1050
 Glu Ile Ala Met His Ser Phe Asn Cys Gly Gly Glu Phe
 340 345 350
 45 TTC TAC TGT AAC ACA ACA GCA CTG TTT AAT AGT ACC TGG 1089
 Phe Tyr Cys Asn Thr Thr Ala Leu Phe Asn Ser Thr Trp
 355 360
 50 AAT GTT ACT AAA GGG TTG AAT AAC ACT GAA GGA AAT AGC 1128
 Asn Val Thr Lys Gly Leu Asn Asn Thr Glu Gly Asn Ser
 365 370 375
 ACA GGA GAT GAA AAT ATC ATA CTC CCA TGT AGA ATA AAA 1167
 Thr Gly Asp Glu Asn Ile Ile Leu Pro Cys Arg Ile Lys
 380 385
 55 CAA ATT ATA AAC ATG TGG CAG GAA GTA GGA AAA GCA ATG 1206
 Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met
 390 395 400
 TAT GCC CCT CCC ATC AGT GGA CAA ATT AGA TGT TCA TCA 1245
 Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser
 405 410 415
 60 AAC ATT ACA GGG CTG CTA CTA ACA AGA GAT GGT GGT AGT 1284
 Asn Ile Thr Gly Leu Leu Thr Arg Asp Gly Gly Ser
 420 425

AAG AAC GAG AGC ATC ACC ACC GAG GTC TTC AGA CCT GGA 1323
 Lys Asn Glu Ser Ile Thr Thr Glu Val Phe Arg Pro Gly
 430 435 440
 5 GGA GGA GAT ATG AGG GAC AAT TGG AGA AGT GAA TTA TAT 1362
 Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr
 445 450
 AAA TAT AAA GTA GTA AAA ATT GAA CCA TTA GGA GTA GCG 1401
 Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala
 455 460 465
 10 CCC ACC AAG GCA AAG AGA AGA GTG GTG CAG AGA GAA AAA 1440
 Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys
 470 475 480
 AGA GCA GTG GGA ACA ATA GGA GCT ATG TTC CTT GGG TTC 1479
 Arg Ala Val Gly Thr Ile Gly Ala Met Phe Leu Gly Phe
 485 490
 15 TTG GGA GCA TAA AGC TTC TAG AGT CGA CCT GCA 1512
 Leu Gly Ala Xaa Ser Phe Xaa Ser Arg Pro Ala
 495 500 504
 20 CLONE C15.3
 CTC GAG GTA CCT GTG TGG AAA GAA GCA ACT ACC ACT 36
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr
 1 5 10
 25 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT AAT ACA GAG 75
 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asn Thr Glu
 15 20 25
 AAA CAT AAT GTT TGG GCC ACA CAC GCC TGT GTA CCC ACA 114
 Lys His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr
 30 35
 30 GAT CCC AAC CCA CAA GAA GTA GTA TTG GGA AAT GTG ACA 153
 Asp Pro Asn Pro Gln Glu Val Val Leu Gly Asn Val Thr
 40 45 50
 GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA 192
 Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln
 55 60
 35 ATG CAT GAA GAT ATA ATC AGT TTA TGG GAT CAA AGT CTA 231
 Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu
 65 70 75
 40 AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT GTT ACT TTA 270
 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu
 80 85 90
 AAT TGC ACT ACT GAT GAT TTA GGG AAT GCT ACT AAT ACC AAT 309
 Asn Cys Thr Asp Asp Leu Gly Asn Ala Thr Asn Thr Asn
 95 100
 45 AGC AGT GCC ACT ACC AAT AGT AGT AGT TGG GAA GAA ATG 348
 Ser Ser Ala Thr Thr Asn Ser Ser Ser Trp Glu Glu Met
 105 110 115
 AAG GGG GAA ATG AAA AGG TGC TCT TTC AAT ATC ACC ACA 387
 Lys Gly Glu Met Lys Arg Cys Ser Phe Asn Ile Thr Thr
 120 125
 50 AGC ATA AGA GAT AAG ATT AAG AAA GAA CAT GCA CTT TTC 426
 Ser Ile Arg Asp Lys Ile Lys Lys Glu His Ala Leu Phe
 130 135 140
 TAT AGA CTT GAT GTA GTA CCA ATA GAT AAT GAT AAT ACC 465
 Tyr Arg Leu Asp Val Val Pro Ile Asp Asn Asp Asn Thr
 145 150 155
 55 ACA TAT AGG TTG ATA AAT TGT AAT ACC TCA GTC ATT ACA 504
 Thr Tyr Arg Leu Ile Asn Cys Asn Thr Ser Val Ile Thr
 160 165
 60 CAG GCC TGT CCA AAG GTA TCA TTT GAG CCA ATT CCC ATA 543
 Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile
 170 175 180
 CAT TTT TGT GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT 582
 His Phe Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys
 185 190

| | | |
|----|---|------|
| | AAT AAT AAG ACG TTC GAG GGA AAA GGA CCA TGT AAA AAT | 621 |
| | Asn Asn Lys Thr Phe Glu Gly Lys Gly Pro Cys Lys Asn | |
| | 195 200 205 | |
| 5 | GTC AGT ACA GTA CAA TGC ACA CAT GGA ATT AGG CCA GTA | 660 |
| | Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val | |
| | 210 215 220 | |
| | GTG TCA ACT CAA CTG CTG TTA AAT GGC AGT CTA GCA GAA | 699 |
| | Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu | |
| | 225 230 | |
| 10 | GAA GAG GTA ATA ATT AGA TCT GGC AAT ATC ACA GAC AAT | 738 |
| | Glu Glu Val Ile Ile Arg Ser Gly Asn Ile Thr Asp Asn | |
| | 235 240 245 | |
| | ACT AAA ACC ATT ATA GTA CAG CTA AAC GAA TCT GTA GTA | 777 |
| | Thr Lys Thr Ile Ile Val Gln Leu Asn Glu Ser Val Val | |
| | 250 255 | |
| 15 | ATT AAT TGT ACA AGA TCC AAC AAC AAT ACA AGA AAA AGT | 816 |
| | Ile Asn Cys Thr Arg Ser Asn Asn Asn Thr Arg Lys Ser | |
| | 260 265 270 | |
| 20 | ATA CAT ATA GGA CCA GGG AGT GCA TTT TTT GCA ACA GGA | 855 |
| | Ile His Ile Gly Pro Gly Ser Ala Phe Phe Ala Thr Gly | |
| | 275 280 285 | |
| | GAA ATA ATA GGA GAT ATA AGA CAA GCA CAC TGT AAC CTT | 894 |
| | Glu Ile Ile Gly Asp Ile Arg Gln Ala His Cys Asn Leu | |
| | 290 295 | |
| 25 | AGT AGA ACA CAA TGG AAT AAC ACT TTA GGA AAG ATA GTC | 933 |
| | Ser Arg Thr Gln Trp Asn Asn Thr Leu Gly Lys Ile Val | |
| | 300 305 310 | |
| | ATA AAA TTA AGA GAA CAA TTT AGA AAA CAA TTT GGA GAA | 972 |
| | Ile Lys Leu Arg Glu Gln Phe Arg Lys Gln Phe Gly Glu | |
| | 315 320 | |
| 30 | AAA ACA ATA GTC TTT AAT CGA TCC TCA GGA GGG GAC CCG | 1011 |
| | Lys Thr Ile Val Phe Asn Arg Ser Ser Gly Gly Asp Pro | |
| | 325 330 335 | |
| | GAA ATT GCA ATG CAC AGT TTT AAT TGT GGA GGG GAA TTT | 1050 |
| | Glu Ile Ala Met His Ser Phe Asn Cys Gly Gly Glu Phe | |
| | 340 345 350 | |
| | TTC TAC TGT AAC ACA ACA GCA CTG TTT AAT AGT ACC TGG | 1089 |
| | Phe Tyr Cys Asn Thr Thr Ala Leu Phe Asn Ser Thr Trp | |
| | 355 360 | |
| 40 | AAT GTT ACT AAA GGG TTG AAT AAC ACT GAA GGA AAT AGC | 1128 |
| | Asn Val Thr Lys Gly Leu Asn Asn Thr Glu Gly Asn Ser | |
| | 365 370 375 | |
| | ACA GGG GAT GAA AAT ATC ATA CTC CCA TGT AGA ATA AAA | 1167 |
| | Thr Gly Asp Glu Asn Ile Ile Leu Pro Cys Arg Ile Lys | |
| | 380 385 | |
| 45 | CAA ATT ATA AAC ATG TGG CAG GAA GTA GGA AAA GCA ATG | 1206 |
| | Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met | |
| | 390 395 400 | |
| | TAT GCC CCT CCC ATC AGT GGA CAA ATT AGA TGT TCA TCA | 1245 |
| | Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser | |
| | 405 410 415 | |
| | AAT ATT ACA GGG CTG CTA CTA ACA AGA GAT GGT GGT AGT | 1284 |
| | Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Ser | |
| | 420 425 | |
| 55 | AAG AAC GAG AGC ATC ACC ACC GAG GTC TTC AGA CCT GGA | 1323 |
| | Lys Asn Glu Ser Ile Thr Thr Glu Val Phe Arg Pro Gly | |
| | 430 435 440 | |
| | GGA GGA GAT ATG AGG GAC AAT TGG AGA AGT GAA TTA TAT | 1362 |
| | Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr | |
| | 445 450 | |
| 60 | AAA TAT AAA GTA GTA AAA ATT GAA CCA TTA GGA GTA GCG | 1401 |
| | Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala | |
| | 455 460 465 | |

CCC ACC AAG GCA AAG AGA AGA GTG GTG CAG AGA GAA AAA 1440
 Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys
 470 475 480
 AGA GCA GTG GGA ACA ATA GGA GCT ATG TTC CTT GGG TTC 1479
 Arg Ala Val Gly Thr Ile Gly Ala Met Phe Leu Gly Phe
 485 490
 TTA GGA GCA TAA AGC TTC TAG A 1501
 Leu Gly Ala Xaa Ser Phe Xaa
 495 500

10

CLONE C7.2

GG GAA TTC GGA TCC GGG GTA CCT GTG TGG AAG GAA GCA 38
 Glu Phe Gly Ser Gly Val Pro Val Trp Lys Glu Ala
 1 5 10
 15 ACC ACC ACT CTA TTC TGT GCA TCA GAT GCT AGA GCA TAT 77
 Thr Thr Thr Leu Phe Cys Ala Ser Asp Ala Arg Ala Tyr
 15 20 25
 GAC ACA GAG GTA CAT AAT GTT TGG GCC ACA CAT GCC TGT 116
 Asp Thr Glu Val His Asn Val Trp Ala Thr His Ala Cys
 30 35
 20 GTA CCC ACA GAC CCT AGT CCA CAA GAA GTA GTT TTG GAA 155
 Val Pro Thr Asp Pro Ser Pro Gln Glu Val Val Leu Glu
 40 45 50
 AAT GTG ACA GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG 194
 Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn Met
 55 60
 25 GTA GAA CAA ATG CAT GAG GAT ATA ATT AGT TTA TGG GAT 233
 Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp
 65 70 75
 30 CAA AGC TTA AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT 272
 Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys
 80 85 90
 GTT ACT TTA AAT TGC AGT GAT TAT AGG AAT GCT ACT GAT 311
 Val Thr Leu Asn Cys Ser Asp Tyr Arg Asn Ala Thr Asp
 95 100
 35 TAT AAG AAT GCT ACT GAT ACC ACT AGT AGT AAC GAG GGA 350
 Tyr Lys Asn Ala Thr Asp Thr Thr Ser Ser Asn Glu Gly
 105 110 115
 40 AAG ATG GAG AGA GGA GAA ATA AAA AAC TGC TCT TTC AAT 389
 Lys Met Glu Arg Gly Glu Ile Lys Asn Cys Ser Phe Asn
 120 125
 ATT ACC ACA AGC ATA AAA AAT AAG ATG CAG AAA GAA TAT 428
 Ile Thr Thr Ser Ile Lys Asn Lys Met Gln Lys Glu Tyr
 130 135 140
 45 GCA CTT TTC TAT AAA CTT GAT ATA GTA CCA ATA GAT AAT 467
 Ala Leu Phe Tyr Lys Leu Asp Ile Val Pro Ile Asp Asn
 145 150 155
 ACA AGC TAT ACA TTG ATA AGT TGT AAC ACC TCA GTC ATT 506
 Thr Ser Tyr Thr Leu Ile Ser Cys Asn Thr Ser Val Ile
 160 165
 50 ACA CAG GCC TGT CCA AAG GTA TCC TTT GAA CCA ACT CCC 545
 Thr Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Thr Pro
 170 175 180
 55 ATA CAT TAT TGT GCT CCG GCT GGT TTT GCG ATT CTA AAG 584
 Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys
 185 190
 TGT AAT GAT AAG AAG TTC AGT GGA AAA GGA GAA TGT AAA 623
 Cys Asn Asp Lys Lys Phe Ser Gly Lys Gly Glu Cys Lys
 195 200 205
 60 AAT GTC AGC ACA GTA CAA TGT ACA CAT GGA ATT AGG CCA 662
 Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro
 210 215 220
 65 GTA GTA TCA ACT CAA CTG CTG TTA AAT GGC AGT CTA GCA 701
 Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala
 225 230

5 GAA GAA GAG GTG GTA ATT AGA TCT GAC AAT TTC ATA GAC 740
 Glu Glu Glu Val Val Ile Arg Ser Asp Asn Phe Ile Asp 235 240 245
 AAT ACT AAA ACC ATA ATA GTA CAG CTG AAA GAA TCT GTA 779
 Asn Thr Lys Thr Ile Ile Val Gln Leu Lys Glu Ser Val 250 255
 10 GAA ATT AAT TGT ATA AGA CCC AAC AAT AAT ACA AGA AAA 818
 Glu Ile Asn Cys Ile Arg Pro Asn Asn Asn Thr Arg Lys 260 265 270
 GGT ATA CAT ATA GGA CCA GGG AGA GCA TGG TAT GCA ACA 857
 Gly Ile His Ile Gly Pro Gly Arg Ala Trp Tyr Ala Thr 275 280 285
 15 GGA GAA ATA GTA GGA GAT ATA AGA AAG GCA TAT TGT AAC 896
 Gly Glu Ile Val Gly Asp Ile Arg Lys Ala Tyr Cys Asn 290 295
 ATT AGT AGA ACA AAA TGG AAT AAC ACT TTA ATA CAG ATA 935
 Ile Ser Arg Thr Lys Trp Asn Asn Thr Leu Ile Gln Ile 300 305 310
 20 GCT AAC AAA TTA AAA GAA AAA TAT AAT ACA ACA ATA ACC 974
 Ala Asn Lys Leu Lys Glu Lys Tyr Asn Thr Thr Ile Ser 315 320
 TTT AAT CGA TCC TCA GGA GGG GAC CCA GAA ATT GTA ACG 1013
 Phe Asn Arg Ser Ser Gly Gly Asp Pro Glu Ile Val Thr 325 330 335
 25 CAT AGT TTT AAT TGT GGA GGG GAG TTT TTC TAC TGT GAT 1052
 His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asp 340 345 350
 TCA ACA CAA CTG TTT AAT AGT ACT TGG AAT TTA AAT GGT 1091
 Ser Thr Gln Leu Phe Asn Ser Thr Trp Asn Leu Asn Gly 355 360
 30 ACT TGG AAT TTT ACT GCA GGG TCA AAT GAA ACT GAA GGC 1130
 Thr Trp Asn Phe Thr Ala Gly Ser Asn Glu Thr Glu Gly 365 370 375
 AAT ATC ACA CTC CCA TGC AGA ATA AAA CAA ATT ATA AAC 1169
 Asn Ile Thr Leu Pro Cys Arg Ile Lys Gln Ile Ile Asn 380 385
 35 AGG TGG CAG GAA GTA GGG AAA GCA ATG TAT GCC CCT CCC 1208
 Arg Trp Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro 390 395 400
 40 ATC AGT GGA CAA ATA AAA TGC TCA TCA AAC ATT ACA GGG 1247
 Ile Ser Gly Gln Ile Lys Cys Ser Ser Asn Ile Thr Gly 405 410 415
 ATG ATA TTA ACA AGG GAT GGT GGT AAC GAG AAC AAT 1286
 Met Ile Leu Thr Arg Asp Gly Gly Asn Glu Asn Asn Asn 420 425
 45 GAG AGC AGT ACT ACT GAG ACC TTC AGA CCG GGA GGA GGA 1325
 Glu Ser Ser Thr Thr Glu Thr Phe Arg Pro Gly Gly Gly 430 435 440
 GAT ATG AGG AAC AAT TGG AGA AGT GAA TTA TAT AAA TAT 1364
 Asp Met Arg Asn Asn Trp Arg Ser Glu Leu Tyr Lys Tyr 445 450
 50 AAA GTA GTA AAA ATT GAA CCA TTA GGA GTA GCA CCC ACC 1403
 Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala Pro Thr 455 460 465
 55 AAG GCA AAG AGA AGA GTG GTG CAG AGA GAA AAA AGA GCA 1442
 Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala 470 475 480
 GTG GGA GCG CTA GGA GCT ATG TTC CTT GGG TTC TTA GGA 1481
 Val Gly Ala Leu Gly Ala Met Phe Leu Gly Phe Leu Gly 485 490
 60 GCA TAA AGC TTC TAG ACC GAC TCT AGA GGA TCC 1514
 Ala Xaa Ser Phe Xaa Thr Asp Ser Arg Gly Ser 495 500 504

CLONE C7.10

| | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| G | GTA | CCT | GTG | TGG | AAG | GAA | GCA | ACC | ACC | ACT | CTA | TTC | 37 |
| | Val | Pro | Val | Trp | Lys | Glu | Ala | Thr | Thr | Thr | Leu | Phe | |
| | 1 | | | | 5 | | | | | 10 | | | |
| 5 | TGT | GCA | TCA | GAT | GCT | AGA | GCA | TAT | GAC | ACA | GAG | GTA | 76 |
| | Cys | Ala | Ser | Asp | Ala | Arg | Ala | Tyr | Asp | Thr | Glu | Val | His |
| | | 15 | | | | | 20 | | | | | 25 | |
| | AAT | GTT | TGG | GCC | ACA | CAT | GCC | TGT | GTA | CCC | ACA | GAC | 115 |
| | Asn | Val | Trp | Ala | Thr | His | Ala | Cys | Val | Pro | Thr | Asp | Pro |
| 10 | | | | | 30 | | | | | 35 | | | |
| | AGT | CCA | CAA | GAA | GTA | TTT | TTG | GGA | AAT | GTG | ACA | GAA | 154 |
| | Ser | Pro | Gln | Glu | Val | Phe | Leu | Gly | Asn | Val | Thr | Glu | Asn |
| | 40 | | | | | | 45 | | | | | 50 | |
| | TTT | AAT | ATG | TGG | AAA | AAT | AAC | ATG | GTA | GAA | CAA | ATG | 193 |
| | Phe | Asn | Met | Trp | Lys | Asn | Asn | Met | Val | Glu | Gln | Met | Tyr |
| 15 | | | | | 55 | | | | 60 | | | | |
| | GAG | GAT | ATA | ATT | AGT | TTA | TGG | GAT | CAA | AGC | TTA | AAG | 232 |
| | Glu | Asp | Ile | Ile | Ser | Leu | Trp | Asp | Gln | Ser | Leu | Lys | Pro |
| | 65 | | | | | 70 | | | | | 75 | | |
| 20 | TGT | GTA | AAA | TTA | ACC | CCA | CTC | TGT | GTT | ACT | TTA | AAT | 271 |
| | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Val | Thr | Leu | Asn | Cys |
| | | | 80 | | | | | 85 | | | | 90 | |
| | AGT | GAT | TAT | AGG | AAT | GCT | ACT | GAT | TAT | AAG | AAT | GCT | 310 |
| | Ser | Asp | Tyr | Arg | Asn | Ala | Thr | Asp | Tyr | Lys | Asn | Ala | Thr |
| 25 | | | | | 95 | | | | | 100 | | | |
| | GAT | ACC | ACT | AGT | AGT | AAC | GAG | GGA | AAG | ATG | GAG | AGA | 349 |
| | Asp | Thr | Thr | Ser | Ser | Asn | Glu | Gly | Lys | Met | Glu | Arg | Gly |
| | 105 | | | | | | 110 | | | | | 115 | |
| | GAA | ATA | AAA | AAC | TGC | TCT | TTC | AAT | ATC | ACC | ACA | AGC | 388 |
| | Glu | Ile | Lys | Asn | Cys | Ser | Phe | Asn | Ile | Thr | Thr | Ser | Ile |
| 30 | | | | | 120 | | | | 125 | | | | |
| | AAA | AAT | AAG | ATG | CAG | AAA | GAA | TAT | GCA | CTT | TTC | TAT | 427 |
| | Lys | Asn | Lys | Met | Gln | Lys | Glu | Tyr | Ala | Leu | Phe | Tyr | Lys |
| | 130 | | | | | 135 | | | | | 140 | | |
| 35 | CTT | AAT | ATA | GTA | CCA | ATA | GAT | AAT | ACA | AGC | TAT | ACA | 466 |
| | Leu | Asn | Ile | Val | Pro | Ile | Asp | Asn | Thr | Ser | Tyr | Thr | Leu |
| | | | 145 | | | | | 150 | | | | 155 | |
| | ATA | AGT | TGT | AAC | ACC | TCA | GTC | ATT | ACA | CAG | GCC | TGT | 505 |
| | Ile | Ser | Cys | Asn | Thr | Ser | Val | Ile | Thr | Gln | Ala | Cys | Pro |
| 40 | | | | | 160 | | | | 165 | | | | |
| | AAG | GTA | TCC | TTT | GAA | CCA | ATT | CCC | ATA | CAT | TAT | TGT | 544 |
| | Lys | Val | Ser | Phe | Glu | Pro | Ile | Pro | Ile | His | Tyr | Cys | Ala |
| | 170 | | | | | | 175 | | | | | 180 | |
| | CCG | GCT | GGT | TTT | GCG | ATT | CTA | AAG | TGT | AAT | GAT | AAG | 583 |
| | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Asn | Asp | Lys | Lys |
| 45 | | | | | 185 | | | | 190 | | | | |
| | TTC | AGT | GGA | AAA | GGA | GAA | TGT | AAA | AAT | GTC | AGC | ACA | 622 |
| | Phe | Ser | Gly | Lys | Gly | Glu | Cys | Lys | Asn | Val | Ser | Thr | Val |
| | 195 | | | | | 200 | | | | 205 | | | |
| 50 | CAA | TGT | ACA | CAT | GGA | ATT | AGG | CCA | GTA | GTA | TCA | ACT | 661 |
| | Gln | Cys | Thr | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | Gln |
| | | | 210 | | | | | 215 | | | | 220 | |
| | CTG | CTG | TTA | AAT | GGC | AGT | CTA | GCA | GAA | GAA | GAG | GTG | 700 |
| | Leu | Leu | Leu | Asn | Gly | Ser | Leu | Ala | Glu | Glu | Glu | Val | Val |
| 55 | | | | | 225 | | | | | 230 | | | |
| | ATT | AGA | TCT | GAC | AAT | TTC | ACA | GAC | AAT | ACT | AAA | ACC | 739 |
| | Ile | Arg | Ser | Asp | Asn | Phe | Thr | Asp | Asn | Thr | Lys | Thr | Ile |
| | 235 | | | | | | 240 | | | | | 245 | |
| | ATA | GTA | CAG | CTG | AAA | GAA | TCT | GTA | GAA | ATT | AAT | TGT | 778 |
| | Ile | Val | Gln | Leu | Lys | Glu | Ser | Val | Glu | Ile | Asn | Cys | Ile |
| 60 | | | | | 250 | | | | 255 | | | | |
| | AGA | CCC | AAC | AAT | AAT | ACA | AGA | AAA | GGT | ATA | CAT | ATA | 817 |
| | Arg | Pro | Asn | Asn | Asn | Thr | Arg | Lys | Gly | Ile | His | Ile | Gly |
| | 260 | | | | | 265 | | | | | 270 | | |

CCA GGG AGA GCA TGG TAT GCA ACA GGA GAA ATA GTA GGA 856
 Pro Gly Arg Ala Trp Tyr Ala Thr Gly Glu Ile Val Gly
 275 280 285
 5 GAT ATA AGA CAG GCA TAT TGT AAC ATT AGT AGA ACA AAA 895
 Asp Ile Arg Gln Ala Tyr Cys Asn Ile Ser Arg Thr Lys
 290 295
 TGG AAT AAC ACT TTA ATA CAG ATA GCT AAC AAA TTA AAA 934
 Trp Asn Asn Thr Leu Ile Gln Ile Ala Asn Lys Leu Lys
 300 305 310
 10 GAA AAA TAT AAT ACA ACA ATA AGC TTT AAT CGA TCC TCA 973
 Glu Lys Tyr Asn Thr Thr Ile Ser Phe Asn Arg Ser Ser
 315 320
 GGA GGG GAC CCA GAA ATT GTA ACC CAT AGT TTT AAT TGT 1012
 Gly Gly Asp Pro Glu Ile Val Thr His Ser Phe Asn Cys
 325 330 335
 15 GGA GGG GAA TTT TTC TAC TGT AAT TCA ACA CAA CTG TTT 1051
 Gly Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe
 340 345 350
 AAT AGT ACT TGG AAT TTA AAT GGT ACT TGG AAT TTT ACT 1090
 Asn Ser Thr Trp Asn Leu Asn Gly Thr Trp Asn Phe Thr
 355 360
 20 GCA GGG TCA AAT GAA ACT GAA GGC AAT ATC ACA CTC CCA 1129
 Ala Gly Ser Asn Glu Thr Glu Gly Asn Ile Thr Leu Pro
 365 370 375
 25 TGC AGA ATA AAA CAA ATT ATA AAC AGG TGG CAG GAA GTA 1158
 Cys Arg Ile Lys Gln Ile Ile Asn Arg Trp Gln Glu Val
 380 385
 GGA AAA GCA ATG TAT GCC CCT CCC ATC AGT GGA CAA ATA 1207
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile
 390 395 400
 30 AGA TGC TCA TCA AAC ATT ACA GGG ATG ATA TTA ACA AGG 1246
 Arg Cys Ser Ser Asn Ile Thr Gly Met Ile Leu Thr Arg
 405 410 415
 GAT GGT GGT AAC GAG AAC AAT AAT GAG AGC AGT ACT ACT 1285
 Asp Gly Gly Asn Glu Asn Asn Asn Glu Ser Ser Thr Thr
 420 425
 35 GAG ACC TTC AGA CCG GGA GGA GGA GAT ATG AGG AAC AAT 1324
 Glu Thr Phe Arg Pro Gly Gly Gly Asp Met Arg Asn Asn
 430 435 440
 40 TGG AGA AGT GAA TTA TAT AAA TAT AAA GTA GTA AAA ATT 1363
 Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 445 450
 GAG CCA TTA GGA GTA GCA CCC ACC GAC TCT AGA GGA TCC 1402
 Glu Pro Leu Gly Val Ala Pro Thr Asp Ser Arg Gly Ser
 455 460 465
 45 TCT AGA 1408
 Ser Arg
 469
 50 CLONE C11.5
 GAG GTA CCT GTG TGG AAA GAA GCA ACC ACT ACT CTA 36
 Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu
 1 5 10
 55 TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GGG GTG 75
 Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Gly Val
 15 20 25
 CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA GAC 114
 His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp
 30 35
 60 CCC AAC CCA CAA GAA ATA GAA TTG GTA AAT GTG ACA GAA 153
 Pro Asn Pro Gln Glu Ile Glu Leu Val Asn Val Thr Glu
 40 45 50
 GAT TTT AAC ATG TGG AAA AAT AAA ATG GTA GAC CAG ATG 192
 Asp Phe Asn Met Trp Lys Asn Lys Met Val Asp Gln Met
 55 60
 65

| | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|------|
| | CAT | GAG | GAT | ATA | ATC | AGT | TTA | TGG | GAT | GAA | AGC | CTA | AAG | 231 |
| | His | Glu | Asp | Ile | Ile | Ser | Leu | Trp | Asp | Glu | Ser | Leu | Lys | |
| | 65 | | | | | 70 | | | | | 75 | | | |
| 5 | CCA | TGT | GTA | AAG | TTA | ACC | CCA | CTT | TGT | GTT | ACT | CTA | AAC | 270 |
| | Pro | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Val | Thr | Leu | Asn | |
| | 80 | | | | | | 85 | | | | | 90 | | |
| | TGC | AGT | GAT | GTG | AAC | AAT | TCC | ACA | AAT | CCT | AAT | GAT | ACT | 309 |
| | Cys | Ser | Asp | Val | Asn | Asn | Ser | Thr | Asn | Pro | Asn | Asp | Thr | |
| | 95 | | | | | | | | 100 | | | | | |
| 10 | AAT | ACT | AAT | TCC | ACT | AAT | ACT | ACT | TCC | TCT | ACT | CCT | ACG | 348 |
| | Asn | Thr | Asn | Ser | Thr | Asn | Thr | Thr | Ser | Ser | Thr | Pro | Thr | |
| | 105 | | | | | | 110 | | | | | 115 | | |
| | GCC | ACT | ACT | AGT | AGC | GAG | GAA | AAG | ATG | GAG | AAG | GGA | GAA | 387 |
| | Ala | Thr | Thr | Ser | Ser | Glu | Glu | Lys | Met | Glu | Lys | Gly | Glu | |
| 15 | | | | 120 | | | | | 125 | | | | | |
| | ATA | AAA | AAC | TGC | TCT | TTC | AAT | ATC | ACC | ACA | CAC | ATG | AAA | 426 |
| | Ile | Lys | Asn | Cys | Ser | Phe | Asn | Ile | Thr | Thr | His | Met | Lys | |
| | 130 | | | | | 135 | | | | | 140 | | | |
| | GAT | AAG | GCA | CAG | AAA | GAA | TAT | GCA | CTT | TTT | TAT | AAA | CTT | 465 |
| 20 | Asp | Lys | Ala | Gln | Lys | Glu | Tyr | Ala | Leu | Phe | Tyr | Lys | Leu | |
| | 145 | | | | | | 150 | | | | | 155 | | |
| | GAT | ATA | GTA | CCA | ATA | GAT | GAT | AAT | AAT | GCC | AGC | TAT | AGG | 504 |
| | Asp | Ile | Val | Pro | Ile | Asp | Asp | Asn | Asn | Ala | Ser | Tyr | Arg | |
| | 160 | | | | | | | | 165 | | | | | |
| 25 | TTG | ATA | AGT | TGT | AAT | ACC | TCA | GAC | ATT | ACA | CAG | GCC | TGT | 543 |
| | Leu | Ile | Ser | Cys | Asn | Thr | Ser | Asp | Ile | Thr | Gln | Ala | Cys | |
| | 170 | | | | | | 175 | | | | | 180 | | |
| | CCA | AAG | GTG | ACC | TTT | GAG | CCA | ATT | CCC | ATA | CAT | TAT | TGT | 582 |
| | Pro | Lys | Val | Thr | Phe | Glu | Pro | Ile | Pro | Ile | His | Tyr | Cys | |
| 30 | | | | 185 | | | | | 190 | | | | | |
| | GCC | CCG | GCT | GGT | TTT | GCG | ATT | CTA | AAG | TGT | AAA | GAT | AAG | 621 |
| | Ala | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Lys | Asp | Lys | |
| | 195 | | | | | 200 | | | | 205 | | | | |
| | AAG | TTC | AAT | GGA | ACA | GGA | CCA | TGT | TCA | AAG | GTC | AGC | ACA | 660 |
| 35 | Lys | Phe | Asn | Gly | Thr | Gly | Pro | Cys | Ser | Lys | Val | Ser | Thr | |
| | 210 | | | | | | 215 | | | | | 220 | | |
| | GTA | CAA | TGT | ACA | CAT | GGA | ATT | AGG | CCA | GTA | GTA | TCA | ACT | 699 |
| | Val | Gln | Cys | Thr | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | |
| | 225 | | | | | | | | 230 | | | | | |
| 40 | CAA | CTG | TTG | TTA | AAT | GGC | AGT | CTT | GCA | GAA | GAA | GTA | 738 | |
| | Gln | Leu | Leu | Leu | Asn | Gly | Ser | Leu | Ala | Glu | Glu | Glu | Val | |
| | 235 | | | | | | 240 | | | | 245 | | | |
| | GTA | ATT | AGA | TCT | GTC | AAT | TTC | ACA | GAC | AAT | GCT | AAA | ATC | 777 |
| 45 | Val | Ile | Arg | Ser | Val | Asn | Phe | Thr | Asp | Asn | Ala | Lys | Ile | |
| | 250 | | | | | | | 255 | | | | | | |
| | ATA | ATA | GTA | CAG | CTG | AAA | GAA | CCT | GTA | GCA | ATT | AAT | TGT | 816 |
| | Ile | Ile | Val | Gln | Leu | Lys | Glu | Pro | Val | Ala | Ile | Asn | Cys | |
| | 260 | | | | | 265 | | | | 270 | | | | |
| | ACA | AGA | CCC | AAC | AAC | AAT | ACA | AGA | AAA | GGT | ATA | CAT | CTA | 855 |
| 50 | Thr | Arg | Pro | Asn | Asn | Asn | Thr | Arg | Lys | Gly | Ile | His | Leu | |
| | 275 | | | | | | 280 | | | | | 285 | | |
| | GGA | CCA | GGG | AGC | ACA | TTT | TAT | ACA | ACA | GGA | GAA | ATA | ATA | 894 |
| | Gly | Pro | Gly | Ser | Thr | Phe | Tyr | Thr | Thr | Gly | Glu | Ile | Ile | |
| | 290 | | | | | | | 295 | | | | | | |
| 55 | GGA | GAC | ATA | AGA | AAA | GCA | TAT | TGC | AAG | ATT | AGT | AAA | GAA | 933 |
| | Gly | Asp | Ile | Arg | Lys | Ala | Tyr | Cys | Lys | Ile | Ser | Lys | Glu | |
| | 300 | | | | | | 305 | | | | | 310 | | |
| | AAA | TGG | AAT | AAC | ACT | TTA | AGA | CAG | GTA | GTT | AAA | AAA | TTA | 972 |
| | Lys | Trp | Asn | Asn | Thr | Leu | Arg | Gln | Val | Val | Lys | Lys | Leu | |
| 60 | | | | 315 | | | | | 320 | | | | | |
| | AGA | GAA | CAA | TTT | GGG | AAT | AAA | ACA | ATA | ATT | TTT | AAT | CGA | 1011 |
| | Arg | Glu | Gln | Phe | Gly | Asn | Lys | Thr | Ile | Ile | Phe | Asn | Arg | |
| | 325 | | | | | 330 | | | | | 335 | | | |

TCC TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC AGT TTT 1050
 Ser Ser Gly Gly Asp Pro Glu Ile Val Met His S r Phe
 340 345 350
 5 AAC TGT GGA GGG GAG TTT TTC TAC TGT AAT ACA ACA CAA 1089
 Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr Gln
 355 360
 CTG TTT AAT AGT ACT TGG AAT AAT ACT GAA GGG ACA AAT 1128
 Leu Phe Asn Ser Thr Trp Asn Asn Thr Glu Gly Thr Asn
 365 370 375
 10 AGC ACT GAA GGA AAT AGC ACA ATC ACA CTC CCA TGC AGA 1167
 Ser Thr Glu Gly Asn Ser Thr Ile Thr Leu Pro Cys Arg
 380 385
 ATA AAA CAA ATT ATA AAT ATG TGG CAG GAA GTA GGA AAA 1206
 Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 390 395 400
 15 GCA ACG TAT GCC CCT CCC ATC AGA GGA CGA ATT AGA TGC 1245
 Ala Thr Tyr Ala Pro Pro Ile Arg Gly Arg Ile Arg Cys
 405 410 415
 20 ATA TCA AAT ATT ACA GGA CTG CTA TTA ACA AGA GAT GGT 1284
 Ile Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly
 420 425
 GGT AGG AAT GTC ACA AAC AAT ACC GAA ACC TTC AGA CCT 1323
 Gly Arg Asn Val Thr Asn Asn Thr Glu Thr Phe Arg Pro
 430 435 440
 25 GGA GGA GGA GAC ATG AGG GAC AAT TGG AGA AGT GAA TTA 1362
 Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu
 445 450
 TAT AAA TAT AAA GTA GTA AAA GTT GAA CCA TTA GGA ATA 1401
 Tyr Lys Tyr Lys Val Val Lys Val Glu Pro Leu Gly Ile
 455 460 465
 30 GCA CCC ACC AAG GCA AAG AGA AGA GTG GTG CAC AGA GAC 1440
 Ala Pro Thr Lys Ala Lys Arg Arg Val Val His Arg Asp
 470 475 480
 35 AAA AGA GCA GCA CTA GGA GCC TTG TTC CTT GGG TTC TTA 1479
 Lys Arg Ala Ala Leu Gly Ala Leu Phe Leu Gly Phe Leu
 485 490
 GGA GCA TAA AAG CTT CTA GA 1499
 Gly Ala Xaa Lys Leu Leu
 495 499
 40
 CLONE C11.7
 GAG GTA CCT GTA TGG AAA GAA GCA ACC ACT ACT CTA 36
 Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu
 1 5 10
 45 TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GAG GTG 75
 Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val
 15 20 25
 CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA GAC 114
 His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp
 30 35
 50 CCC AAC CCA CAA GAA ATA GAA TTG GTA AAT GTG ACA GAA 153
 Pro Asn Pro Gln Glu Ile Glu Leu Val Asn Val Thr Glu
 40 45 50
 55 GAT TTT AAC ATG TGG AAA AAT AAA ATG GTA GAC CAG ATG 192
 Asp Phe Asn Met Trp Lys Asn Lys Met Val Asp Gln Met
 55 60
 CAT GAG GAT ATA ATC AGT TTA TGG GAT GAA AGC CTA AAG 231
 His Glu Asp Ile Ile Ser Leu Trp Asp Glu Ser Leu Lys
 65 70 75
 60 CCA TGT GTA AAG TTA ACC CCA CTT TGT GTT ACT CTA AAC 270
 Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn
 80 85 90
 TGC AGT GAT GTG AAC AAT TCC ACA AAT CCT AAT GAT ACT 309
 Cys Ser Asp Val Asn Asn Ser Thr Asn Pro Asn Asp Thr
 95 100
 65

| | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|------|
| | AAT | ACT | AAT | TCC | ACT | AAT | ACT | ACT | TCC | TCT | ACT | CCT | ACG | 348 |
| | Asn | Thr | Asn | Ser | Thr | Asn | Thr | Thr | Ser | Ser | Thr | Pro | Thr | |
| | 105 | | | | | 110 | | | | | | 115 | | |
| 5 | GCC | ACT | ACT | AGT | AGC | GAG | GAA | AAG | ATG | GAG | AAG | GGA | GAA | 387 |
| | Ala | Thr | Thr | Ser | Ser | Glu | Glu | Lys | Met | Glu | Lys | Gly | Glu | |
| | 120 | | | | | 125 | | | | | | | | |
| | ATA | AAA | AAC | TGC | TCT | TTC | AAT | ATC | ACC | ACA | CAC | ATG | AAA | 426 |
| | Ile | Lys | Asn | Cys | Ser | Phe | Asn | Ile | Thr | Thr | His | Met | Lys | |
| | 130 | | | | | 135 | | | | | 140 | | | |
| 10 | GAT | AAG | GTA | CAG | AAA | GAA | TAT | GCA | CTT | TTT | TAT | AAA | CTT | 465 |
| | Asp | Lys | Val | Gln | Lys | Glu | Tyr | Ala | Leu | Phe | Tyr | Lys | Leu | |
| | 145 | | | | | 150 | | | | | | | 155 | |
| | GAT | ATA | GTA | CCA | ATA | GAT | GAT | AAT | AAT | ACC | AGC | TAT | AGG | 504 |
| | Asp | Ile | Val | Pro | Ile | Asp | Asp | Asn | Asn | Thr | Ser | Tyr | Arg | |
| | 160 | | | | | 165 | | | | | | | | |
| 15 | TTG | ATA | AGT | TGT | AAT | ACC | TCA | GTC | ATT | ACA | CAG | GCC | TGT | 543 |
| | Leu | Ile | Ser | Cys | Asn | Thr | Ser | Val | Ile | Thr | Gln | Ala | Cys | |
| | 170 | | | | | 175 | | | | | | 180 | | |
| 20 | CCA | ATG | GTG | ACC | TTT | GAG | CCA | ATT | CCC | ATA | CAT | TAT | TGT | 582 |
| | Pro | Met | Val | Thr | Phe | Glu | Pro | Ile | Pro | Ile | His | Tyr | Cys | |
| | 185 | | | | | 190 | | | | | | | | |
| | GCC | CCG | GCT | GGT | TTT | GCG | ATT | CTA | AAG | TGT | AAA | GAT | AAG | 621 |
| | Ala | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Lys | Asp | Lys | |
| | 195 | | | | | 200 | | | | | 205 | | | |
| 25 | AAG | TTC | AAT | GGA | ACA | GGA | CCA | TGT | TCA | AAG | GTC | AGC | ACA | 660 |
| | Lys | Phe | Asn | Gly | Thr | Gly | Pro | Cys | Ser | Lys | Val | Ser | Thr | |
| | 210 | | | | | 215 | | | | | | | 220 | |
| | GTA | CAA | TGT | ACA | CAT | GGA | ATT | AGG | CCA | GTA | GTA | TCA | ACT | 699 |
| | Val | Gln | Cys | Thr | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | |
| | 225 | | | | | 230 | | | | | | | | |
| 30 | CAA | CTG | TTG | TTA | AAT | GGC | AGT | CTT | GCA | GAA | GAA | GAA | GTA | 738 |
| | Gln | Leu | Leu | Leu | Asn | Gly | Ser | Leu | Ala | Glu | Glu | Glu | Val | |
| | 235 | | | | | 240 | | | | | | 245 | | |
| 35 | GTA | ATT | AGA | TCT | GTC | AAT | TTC | ACA | GAC | AAT | GCT | AAA | ATC | 777 |
| | Val | Ile | Arg | Ser | Val | Asn | Phe | Thr | Asp | Asn | Ala | Lys | Ile | |
| | 250 | | | | | 255 | | | | | | | | |
| | ATA | ATA | GTA | CAG | CTG | AAA | GAA | CCT | GTA | GCA | ATT | AAT | TGT | 816 |
| | Ile | Ile | Val | Gln | Leu | Lys | Glu | Pro | Val | Ala | Ile | Asn | Cys | |
| | 260 | | | | | 265 | | | | | 270 | | | |
| 40 | ACA | AGA | CCC | AAC | AAC | AAT | ACA | AGA | AAA | GGT | ATA | CAT | CTA | 855 |
| | Thr | Arg | Pro | Asn | Asn | Asn | Thr | Arg | Lys | Gly | Ile | His | Leu | |
| | 275 | | | | | 280 | | | | | | | 285 | |
| | GGA | CCA | GGG | AGC | ACA | TTT | TAT | ACA | ACA | GGA | GAA | ATA | ATA | 894 |
| | Gly | Pro | Gly | Ser | Thr | Phe | Tyr | Thr | Thr | Gly | Glu | Ile | Ile | |
| | 290 | | | | | 295 | | | | | | | | |
| 45 | GGA | GAC | ATA | AGA | AAA | GCA | TAT | TGC | AAG | ATT | AGT | AAA | GAA | 933 |
| | Gly | Asp | Ile | Arg | Lys | Ala | Tyr | Cys | Lys | Ile | Ser | Lys | Glu | |
| | 300 | | | | | 305 | | | | | | 310 | | |
| 50 | AAA | TGG | AAT | AAC | ACT | TTA | AGA | CAG | GTA | GTT | AAA | AAA | TTA | 972 |
| | Lys | Trp | Asn | Asn | Thr | Leu | Arg | Gln | Val | Val | Lys | Lys | Leu | |
| | 315 | | | | | 320 | | | | | | | | |
| | AGA | GAA | CAA | TTT | GGG | AAT | AAA | ACA | ATA | ATT | TTT | AAT | CGA | 1011 |
| | Arg | Glu | Gln | Phe | Gly | Asn | Lys | Thr | Ile | Ile | Phe | Asn | Arg | |
| | 325 | | | | | 330 | | | | | 335 | | | |
| 55 | TCC | TCA | GGA | GGG | GAC | CCA | GAA | ATT | GTA | ATG | CAC | AGT | TTT | 1050 |
| | Ser | Ser | Gly | Gly | Asp | Pro | Glu | Ile | Val | Met | His | Ser | Phe | |
| | 340 | | | | | 345 | | | | | | | 350 | |
| | AAC | TGT | GGA | GGG | GAG | TTT | TTC | TAC | TGT | AAT | ACA | ACA | CAA | 1089 |
| | Asn | Cys | Gly | Gly | Glu | Phe | Phe | Tyr | Cys | Asn | Thr | Thr | Gln | |
| | 355 | | | | | 360 | | | | | | | | |
| 60 | CTG | TTT | AAT | AGT | ACT | TGG | AAT | AAT | ACT | GAA | GGG | ACA | AAT | 1128 |
| | Leu | Phe | Asn | Ser | Thr | Trp | Asn | Asn | Thr | Glu | Gly | Thr | Asn | |
| | 365 | | | | | 370 | | | | | | 375 | | |

AGC ACT GAA GGA AAT AGC ACA ATC ACA CTC CCA TGC AGA 1167
 Ser Thr Glu Gly Asn Ser Thr Il Thr Leu Pro Cys Arg
 380 385
 5 ATA AAA CAA ATT ATA AAT ATG TGG CAG GAA GTA GGA AAA 1206
 Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 390 395 400
 GCA ACG TAT GCC CCT CCC ATC AGA GGA CGA ATT AGA TGC 1245
 Ala Thr Tyr Ala Pro Pro Ile Arg Gly Arg Ile Arg Cys
 405 410 415
 10 ATA TCA AAT ATT ACA GGA CTG CTA TTA ACA AGA GAT GGT 1284
 Ile Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly
 420 425
 GGT AGG AAT GTC ACA AAC AAT ACC GAN NCC TTC AGA CCT 1323
 Gly Arg Asn Val Thr Asn Asn Thr Xaa Xaa Phe Arg Pro
 430 435 440
 15 GGA GGA GGA GAC ATG AGG GAC AAT TGG AGA AGT GAA TTA 1362
 Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu
 445 450
 TAT AAA TAT AAA GTA GTA AAA GTT GAA CCA TTA GGA ATA 1401
 Tyr Lys Tyr Lys Val Val Lys Val Glu Pro Leu Gly Ile
 455 460 465
 20 GCA CCC ACC AAG GCA AAG AGA AGA GTG GTG CAC AGA GAC 1440
 Ala Pro Thr Lys Ala Lys Arg Arg Val Val His Arg Asp
 470 475 480
 25 AAA AGA GCA GCA CTA GGA GCT TTG TTC CTT GGC TTC TTA 1479
 Lys Arg Ala Ala Leu Gly Ala Leu Phe Leu Gly Phe Leu
 485 490
 GGA GCA TAA AAG CTT CTA GA 1499
 Gly Ala Xaa Lys Leu Leu
 495 499
 30

CLONE C10.5

6 GTA CCT GTG TGG AAA GAA GCA AAC ACA ACT CTA TTT 37
 Val Pro Val Trp Lys Glu Ala Asn Thr Thr Leu Phe
 1 5 10
 35 TGT GCA TCA GAT GCT AAA GCA TAT GAT AGA GAA GTA CAT 76
 Cys Ala Ser Asp Ala Lys Ala Tyr Asp Arg Glu Val His
 15 20 25
 40 AAT GTT TGG GCA ACA CAT GCC TGT GTA CCC ACA GAC CCC 115
 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro
 30 35
 AAC CCA CAA GAA ATA GTA TTG GGA AAT GTG ACA GAA AAT 154
 Asn Pro Gln Glu Ile Val Leu Gly Asn Val Thr Glu Asn
 40 45 50
 45 TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA ATG CAT 193
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His
 55 60
 GAG GAT ATA ATC AAT TTA TGG GAT CAA AGC TTA AAG CCA 232
 Glu Asp Ile Ile Asn Leu Trp Asp Gln Ser Leu Lys Pro
 65 70 75
 50 TGT GTA AAG TTA ACT CCA CTC TGT GTT ACT TTA AAG TGC 271
 Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Lys Cys
 80 85 90
 AAG GAT CTG GAG AGG AAT ACT ACC TAT AAT AGC ACT ATT 310
 Lys Asp Leu Glu Arg Asn Thr Thr Tyr Asn Ser Thr Ile
 95 100
 ACC AAT AAT AGT AGT TTG GAG GGA CTA AGA GAA CAA ATG 349
 Thr Asn Asn Ser Ser Leu Glu Gly Leu Arg Glu Gln Met
 105 110 115
 60 ACA AAC TGC TCT TTC AAC ATC ACC ACA AGT ATA AGA GAT 388
 Thr Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Arg Asp
 120 125
 AAG GTG CAG AAA GAA TAT GCA CTT TTG TAT AAA CTT GAT 427
 Lys Val Gln Lys Glu Tyr Ala Leu Leu Tyr Lys Leu Asp
 130 135 140
 65

| | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|------|
| | GTA | GTA | CCA | ATA | GAA | GAA | GAT | GAC | AAT | ACT | AGC | TAT | AGA | 466 |
| | Val | Val | Pro | Ile | Glu | Glu | Asp | Asp | Asn | Thr | Ser | Tyr | Arg | |
| | | | 145 | | | | | 150 | | | | | 155 | |
| 5 | TTG | ATA | AGT | TGT | AAC | ACC | TCA | GTC | ATT | ACA | CAG | GCT | TGT | 505 |
| | Leu | Ile | Ser | Cys | Asn | Thr | Ser | Val | Ile | Thr | Gln | Ala | Cys | |
| | | | | 160 | | | | | | | 165 | | | |
| | CCA | AAG | ACA | TCC | TTT | GAG | CCA | ATT | CCC | ATA | CAT | TAT | TGT | 544 |
| | Pro | Lys | Thr | Ser | Phe | Glu | Pro | Ile | Pro | Ile | His | Tyr | Cys | |
| | | 170 | | | | | 175 | | | | | 180 | | |
| 10 | GCC | CCG | GCT | GGT | TTT | GCG | ATT | CTA | AAG | TGT | AAT | GAT | AAG | 583 |
| | Ala | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Asn | Asp | Lys | |
| | | | | 185 | | | | | 190 | | | | | |
| | AAG | TTC | AAT | GGA | ACA | GGA | CCA | TGT | AAA | AAT | GTC | AGC | ACA | 622 |
| | Lys | Phe | Asn | Gly | Thr | Gly | Pro | Cys | Lys | Asn | Val | Ser | Thr | |
| 15 | | 195 | | | | 200 | | | | | 205 | | | |
| | GTA | CAA | TGT | ACA | CAT | GGA | ATT | AGG | CCA | GTA | GTA | TCA | ACT | 661 |
| | Val | Gln | Cys | Thr | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | |
| | | 210 | | | | | 215 | | | | | 220 | | |
| 20 | CAA | CTG | TTG | TTA | AAT | GCG | AGT | CTA | GCA | GAA | GAA | GAG | GTA | 700 |
| | Gln | Leu | Leu | Leu | Asn | Gly | Ser | Leu | Ala | Glu | Glu | Glu | Val | |
| | | | | 225 | | | | | | 230 | | | | |
| | GTA | ATC | AGA | TCT | GCC | AAT | TTC | ACA | GAC | AAT | GCT | AAA | ACC | 739 |
| | Val | Ile | Arg | Ser | Ala | Asn | Phe | Thr | Asp | Asn | Ala | Lys | Thr | |
| | | 235 | | | | | 240 | | | | | 245 | | |
| 25 | ATA | ATA | GTA | CAT | CTA | AAT | GAA | ACT | GTA | AAA | ATT | AAT | TGT | 778 |
| | Ile | Ile | Val | His | Leu | Asn | Glu | Thr | Val | Lys | Ile | Asn | Cys | |
| | | | | 250 | | | | 255 | | | | | | |
| | ACA | AGA | CTT | GGC | AAC | AAT | ACA | AGA | AAA | AGT | ATA | AAT | ATA | 817 |
| | Thr | Arg | Leu | Gly | Asn | Asn | Thr | Arg | Lys | Ser | Ile | Asn | Ile | |
| 30 | | 260 | | | | 265 | | | | | 270 | | | |
| | GGA | CCA | GGG | AGA | GTA | CTC | TAT | GCA | ACA | GGA | GAA | ATA | ATA | 856 |
| | Gly | Pro | Gly | Arg | Val | Leu | Tyr | Ala | Thr | Gly | Glu | Ile | Ile | |
| | | | 275 | | | | 280 | | | | | 285 | | |
| 35 | GGA | GAC | ATA | AGA | CAA | GCA | CAT | TGT | AAC | ATT | AGT | AGA | GCA | 895 |
| | Gly | Asp | Ile | Arg | Gln | Ala | His | Cys | Asn | Ile | Ser | Arg | Ala | |
| | | | | 290 | | | | | | 295 | | | | |
| | CAA | TGG | AAT | AAG | ACT | TTA | GAA | AAG | GTA | GTT | GAC | AAA | TTA | 934 |
| | Gln | Trp | Asn | Lys | Thr | Leu | Glu | Lys | Val | Val | Asp | Lys | Leu | |
| | | 300 | | | | | 305 | | | | | 310 | | |
| 40 | AGA | AAA | CAA | TTT | GGG | GAT | AAT | ACA | ACA | ATA | GCT | TTT | AAT | 973 |
| | Arg | Lys | Gln | Phe | Gly | Asp | Asn | Thr | Thr | Ile | Ala | Phe | Asn | |
| | | | 315 | | | | | 320 | | | | | | |
| | CGA | TCC | TCA | GGG | GGG | GAC | CCA | GAA | ATT | GTA | ATG | CAC | ACT | 1012 |
| | Arg | Ser | Ser | Gly | Gly | Asp | Pro | Glu | Ile | Val | Met | His | Thr | |
| 45 | | 325 | | | | 330 | | | | | 335 | | | |
| | TTT | AAT | TGT | GGA | GGG | GAA | TTT | TTC | TAC | TGT | AAT | ACA | ACA | 1051 |
| | Phe | Asn | Cys | Gly | Gly | Glu | Phe | Phe | Tyr | Cys | Asn | Thr | Thr | |
| | | | 340 | | | | 345 | | | | | 350 | | |
| 50 | CAA | CTG | TTT | AAT | AGT | ACT | TGG | AAT | AAT | ACT | TGG | AAG | GAT | 1090 |
| | Gln | Leu | Phe | Asn | Ser | Thr | Trp | Asn | Asn | Thr | Trp | Lys | Asp | |
| | | | | 355 | | | | | | 360 | | | | |
| | CCT | AAC | AGG | AGT | GAC | AAT | ATC | ACA | CTC | CCA | TGC | AGA | ATA | 1129 |
| | Pro | Asn | Arg | Ser | Asp | Asn | Ile | Thr | Leu | Pro | Cys | Arg | Ile | |
| | | 365 | | | | | 370 | | | | | 375 | | |
| 55 | AAA | CAA | ATT | ATA | AAC | ATG | TGG | CAG | GAA | GTA | GGA | AAA | GCA | 1168 |
| | Lys | Gln | Ile | Ile | Asn | Met | Trp | Gln | Glu | Val | Gly | Lys | Ala | |
| | | | | 380 | | | | 385 | | | | | | |
| | ATG | TAC | GCC | CCT | CCC | ATC | AGA | GGG | GAA | ATT | AGA | TGT | TCA | 1207 |
| | Met | Tyr | Ala | Pro | Pro | Ile | Arg | Gly | Glu | Ile | Arg | Cys | Ser | |
| | | 390 | | | | 395 | | | | | 400 | | | |
| 60 | TCA | AAT | ATC | ACA | GGG | CTG | CTA | CTA | ACA | AGA | GAT | GGT | GGT | 1246 |
| | Ser | Asn | Ile | Thr | Gly | Leu | Leu | Leu | Thr | Arg | Asp | Gly | Gly | |
| | | | | 405 | | | 410 | | | | | 415 | | |

AAT GAC GAT GGT AAT GAC ACG ACC ACA AAC AGG ACC GAG 1285
 Asn Asp Asp Gly Asn Asp Thr Thr Thr Asn Arg Thr Glu
 420 425
 5 ATC TTC AGA CCT GGA GGA GGA GAT ATG AGG GAC AAT TGG 1324
 Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp
 430 435 440
 AGA AGT GAA TTA TAT AGA TAT AAA GTA GTA AAA ATT GAA 1363
 Arg Ser Glu Leu Tyr Arg Tyr Lys Val Val Lys Ile Glu
 445 450
 10 CCA TTA GGA ATA GCA CCC ACC AGG GCA AAG AGA AGA GTG 1402
 Pro Leu Gly Ile Ala Pro Thr Arg Ala Lys Arg Arg Val
 455 460 465
 GTG CAG AGA GAA AAA AGA GCA GTA GGA CTA GGA GCT TTG 1441
 Val Gln Arg Glu Lys Arg Ala Val Gly Leu Gly Ala Leu
 470 475 480
 15 TTC CTT GGG T TCTTAGGAG CATAAAGCTT CTAGA 1475
 Phe Leu Gly
 483
 20 CLONE C10.7
 G GTA CCT GTG TGG AAA GAA GCA AAC ACA ACT CTA TTT 37
 Val Pro Val Trp Lys Glu Ala Asn Thr Thr Leu Phe
 1 5 10
 25 TGT GCA TCA GAT GCT AAA GCA TAT GAT AGA GAA GTA CAT 76
 Cys Ala Ser Asp Ala Lys Ala Tyr Asp Arg Glu Val His
 15 20 25
 AAT GTT TGG GCA ACA CAT GCC TGT GTA CCC ACA GAC CCC 115
 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro
 30 35
 AAC CCA CAA GAA ATA GTA TTG GGA AAT GTG ACA GAA AAT 154
 Asn Pro Gln Glu Ile Val Leu Gly Asn Val Thr Glu Asn
 40 45 50
 TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA ATG CAT 193
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His
 55 60
 35 GAG GAT ATA ATC AAT TTA TGG GAT CAA AGC TTA AAG CCA 232
 Glu Asp Ile Ile Asn Leu Trp Asp Gln Ser Leu Lys Pro
 65 70 75
 TGT GTA AAG TTA ACT CCA CTC TGT GTT ACT TTA AAG TGC 271
 Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Lys Cys
 80 85 90
 40 AAG GAT CTG GAG AGG AAT ACT ACC TAT AAT AGC ACT ATT 310
 Lys Asp Leu Glu Arg Asn Thr Thr Tyr Asn Ser Thr Ile
 95 100
 45 ACC AAT AAT AGT AGT TTG GAG GGA CTA AGA GAA CAA ATG 349
 Thr Asn Asn Ser Ser Leu Glu Gly Leu Arg Glu Gln Met
 105 110 115
 ACA AAC TGC TCT TTC AAC ATC ACC ACA AGT ATA AGA GAT 388
 Thr Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Arg Asp
 120 125
 50 AAG GTG CAG AAA GAA TAT GCA CTT TTG TAT AAA CTT GAT 427
 Lys Val Gln Lys Glu Tyr Ala Leu Leu Tyr Lys Leu Asp
 130 135 140
 GTA GTA CCA ATA GAA GAA GAT GAC AAT ACT AGC TAT AGA 466
 Val Val Pro Ile Glu Glu Asp Asp Asn Thr Ser Tyr Arg
 145 150 155
 55 TTG ATA AGT TGT AAC ACC TCA GTC ATT ACA CAG GCT TGT 505
 Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys
 160 165
 60 CCA AAG ACA TCC TTT GAG CCA ATT CCC ATA CAT TAT TGT 544
 Pro Lys Thr Ser Phe Glu Pro Ile Pro Ile His Tyr Cys
 170 175 180
 GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT AAT GAT AAG 583
 Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asp Lys
 185 190

| | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|------|
| | AAG | TTC | AAT | GGA | ACA | GGA | CCA | TGT | AAA | AAT | GTC | AGC | ACA | 622 |
| | Lys | Phe | Asn | Gly | Thr | Gly | Pro | Cys | Lys | Asn | Val | Ser | Thr | |
| | 195 | | | | | 200 | | | | | 205 | | | |
| 5 | GTA | CAA | TGT | ACA | CAT | GGA | ATT | AGG | CCA | GTA | GTA | TCA | ACT | 661 |
| | Val | Gln | Cys | Thr | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | |
| | | | 210 | | | | | 215 | | | | | 220 | |
| | CAA | CTG | TTG | TTA | AAT | GGC | AGT | CTA | GCA | GAA | GAA | GAG | GTA | 700 |
| | Gln | Leu | Leu | Leu | Asn | Gly | Ser | Leu | Ala | Glu | Glu | Glu | Val | |
| | | | | | 225 | | | | | 230 | | | | |
| 10 | GTA | ATC | AGA | TCT | GCC | AAT | TTC | ACA | GAC | AAT | GCT | AAA | ACC | 739 |
| | Val | Ile | Arg | Ser | Ala | Asn | Phe | Thr | Asp | Asn | Ala | Lys | Thr | |
| | | 235 | | | | | 240 | | | | | 245 | | |
| | ATA | ATA | GTA | CAT | CTA | AAT | GAA | ACT | GTA | AAA | ATT | AAT | TGT | 778 |
| | Ile | Ile | Val | His | Leu | Asn | Glu | Thr | Val | Lys | Ile | Asn | Cys | |
| 15 | | | | | 250 | | | | | 255 | | | | |
| | ACA | AGA | CTT | GGC | AAC | AAT | ACA | AGA | AAA | AGT | ATA | AAT | ATA | 817 |
| | Thr | Arg | Leu | Gly | Asn | Asn | Thr | Arg | Lys | Ser | Ile | Asn | Ile | |
| | 260 | | | | | 265 | | | | | 270 | | | |
| 20 | GGA | CCA | GGG | AGA | GTA | CTC | TAT | GCA | ACA | GGA | GAA | ATA | ATA | 856 |
| | Gly | Pro | Gly | Arg | Val | Leu | Tyr | Ala | Thr | Gly | Glu | Ile | Ile | |
| | | | 275 | | | | | 280 | | | | | 285 | |
| | GGA | GAC | ATA | AGA | CAA | GCA | CAT | TGT | AAC | ATT | AGT | AGA | GCA | 895 |
| | Gly | Asp | Ile | Arg | Gln | Ala | His | Cys | Asn | Ile | Ser | Arg | Ala | |
| | | | | | 290 | | | | | 295 | | | | |
| 25 | CAA | TGG | AAT | AAG | ACT | TTA | GAA | AAG | GTA | GTT | GAC | AAG | TTA | 934 |
| | Gln | Trp | Asn | Lys | Thr | Leu | Glu | Lys | Val | Val | Asp | Lys | Leu | |
| | | 300 | | | | | 305 | | | | | 310 | | |
| | AGA | AAA | CAA | TTT | GGG | GAT | AAT | ACA | ACA | ATA | GCT | TTT | AAT | 973 |
| | Arg | Lys | Gln | Phe | Gly | Asp | Asn | Thr | Thr | Ile | Ala | Phe | Asn | |
| 30 | | | | | 315 | | | | | 320 | | | | |
| | CGA | TCC | TCA | GGA | GGG | GAC | CCA | GAA | ATT | GTA | ATG | CAC | ACT | 1012 |
| | Arg | Ser | Ser | Gly | Gly | Asp | Pro | Glu | Ile | Val | Met | His | Thr | |
| | | 325 | | | | 330 | | | | | 335 | | | |
| 35 | TTT | AAT | TGT | GGA | GGG | GAA | TTT | TTC | TAC | TGT | AAT | ACA | ACA | 1051 |
| | Phe | Asn | Cys | Gly | Gly | Glu | Phe | Phe | Tyr | Cys | Asn | Thr | Thr | |
| | | | 340 | | | | | 345 | | | | | 350 | |
| | CAA | CTG | TTT | AAT | AGT | ACT | TGG | AAT | AAT | ACT | TGG | AAG | GAT | 1090 |
| | Gln | Leu | Phe | Asn | Ser | Thr | Trp | Asn | Asn | Thr | Trp | Lys | Asp | |
| | | | | | 355 | | | | | 360 | | | | |
| 40 | CCT | AAC | AGG | AGT | GAC | AAT | ATC | ACA | CTC | CCA | TGC | AGA | ATA | 1129 |
| | Pro | Asn | Arg | Ser | Asp | Asn | Ile | Thr | Leu | Pro | Cys | Arg | Ile | |
| | | 365 | | | | | 370 | | | | | 375 | | |
| | AAA | CAA | ATT | ATA | AAC | ATG | TGG | CAG | GAA | GTA | GGA | AAA | GCA | 1168 |
| | Lys | Gln | Ile | Ile | Asn | Met | Trp | Gln | Glu | Val | Gly | Lys | Ala | |
| 45 | | | | | 380 | | | | | 385 | | | | |
| | ATG | TAC | GCC | CCT | CCC | ATC | AGA | GGG | GAA | ATT | AGA | TGT | TCA | 1207 |
| | Met | Tyr | Ala | Pro | Pro | Ile | Arg | Gly | Glu | Ile | Arg | Cys | Ser | |
| | | 390 | | | | 395 | | | | | 400 | | | |
| | TCA | AAT | ATC | ACA | GGG | CTG | CTA | CTA | ACA | AGA | GAT | GGT | GGT | 1246 |
| | Ser | Asn | Ile | Thr | Gly | Leu | Leu | Leu | Thr | Arg | Asp | Gly | Gly | |
| 50 | | | | | 405 | | | | 410 | | | | 415 | |
| | AAT | GAC | GAT | GGT | AAT | GAC | ACG | ACC | ACA | AAC | AGG | ACC | GAG | 1285 |
| | Asn | Asp | Asp | Gly | Asn | Asp | Thr | Thr | Thr | Asn | Arg | Thr | Glu | |
| | | | | | 420 | | | | | 425 | | | | |
| 55 | ATC | TTC | AGA | CCT | GGA | GGA | GGA | GAT | ATG | AGG | GAC | AAT | TGG | 1324 |
| | Ile | Phe | Arg | Pro | Gly | Gly | Gly | Asp | Met | Arg | Asp | Asn | Trp | |
| | | 430 | | | | | 435 | | | | | 440 | | |
| | AGA | AGT | GAA | TTA | TAT | AGA | TAT | AAA | GTA | GTA | AAA | ATT | GAA | 1363 |
| | Arg | Ser | Glu | Leu | Tyr | Arg | Tyr | Lys | Val | Val | Lys | Ile | Glu | |
| 60 | | | | | 445 | | | | 450 | | | | | |
| | CCA | TTA | GGA | ATA | GCA | CCC | ACC | AGG | GCA | AAG | AGA | AGA | GTG | 1402 |
| | Pro | Leu | Gly | Ile | Ala | Pro | Thr | Arg | Ala | Lys | Arg | Arg | Val | |
| | | 455 | | | | 460 | | | | | 465 | | | |

GTG CAG AGA GAA AAA AGA GCA GTA GGA CTA GGA GCT TTG 1441
 Val Gln Arg Glu Lys Arg Ala Val Gly Leu Gly Ala Leu
 470 475 480
 TTC CTT GGG TTC TTG GGA GCA TAA AGC TTC TAG A 1475
 5 Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa
 485 490 491

CLONE C17.1

10 CTC GAG GTA CCT GTG TGG AAA GAA GCA ACC ACC ACT 36
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr
 1 5 10
 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT GAT TCA GAG 75
 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Ser Glu
 15 15 20 25
 15 GCA CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA 114
 Ala His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr
 30 35
 GAC CCC AAC CCA CAA GAA GTA GAA TTG GAA AAT GTG ACA 153
 Asp Pro Asn Pro Gln Glu Val Glu Leu Glu Asn Val Thr
 20 40 45 50
 GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAG 192
 Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln
 55 60
 25 ATG CAT GGG GAT ATA ATT AGT TTA TGG GAT CAA AGC CTA 231
 Met His Gly Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu
 65 70 75
 AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT GTT ACG TTA 270
 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu
 80 85 90
 30 AAT TGC ACT GAC CCA AAT GTT ACT AAT AGC GAG AGA ACG 309
 Asn Cys Thr Asp Pro Asn Val Thr Asn Ser Glu Arg Thr
 95 100
 ATA GAG GGG GGA GAA ATA AAA AAT TGC TCT TTC AAT ATC 348
 Ile Glu Gly Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile
 35 105 110 115
 ACC ACA AAC ATA AGA GAT AGG TTT CAG AAA GAA TAT GCA 387
 Thr Thr Asn Ile Arg Asp Arg Phe Gln Lys Glu Tyr Ala
 120 125
 40 CTT TTT TAT AAA CTT GAT GTA ATA CCA TTA GGT AAT GAT 426
 Leu Phe Tyr Lys Leu Asp Val Ile Pro Leu Gly Asn Asp
 130 135 140
 AAT ACT AGC TAT AGG TTG ATA AGT TGT AAC ACC TCA GTC 465
 Asn Thr Ser Tyr Arg Leu Ile Ser Cys Asn Thr Ser Val
 145 150 155
 45 ATT ACA CAG GCC TGT CCA AAG GTA TCC TTT GAG CCA ATT 504
 Ile Thr Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile
 160 165
 CCC ATA CAT TAT TGT GCC CCG GCT GGT TTT GCG ATT CTA 543
 Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile Leu
 50 170 175 180
 AAG TGT AAA GAT AAG AAG TTC AAT GGA ACA GGA CCA TGT 582
 Lys Cys Lys Asp Lys Lys Phe Asn Gly Thr Gly Pro Cys
 185 190
 ACA AAT GTC AGC ACA GTA CAA TGT ACA CAT GGA ATT AAG 621
 Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Lys
 55 195 200 205
 CCA GTA GTA TCA ACT CAA CTG TTG TTA AAT GGC AGT CTA 660
 Pro Val Val Ser Thr Gln Leu Leu Asn Gly Ser Leu
 210 215 220
 60 GCA GAA GAA GAC ATA GTA ATT AGA TCC GCC AAT CTC ACA 699
 Ala Glu Glu Asp Ile Val Ile Arg Ser Ala Asn Leu Thr
 225 230
 GAC AAT GCT AAA AAC ATA ATA GTA CAG CTG AAT GAA TCT 738
 Asp Asn Ala Lys Asn Ile Ile Val Gln Leu Asn Glu Ser
 65 235 240 245

-37-

| | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| | GAC | CCC | AAC | CCA | CAA | GAA | GTA | GAA | TTG | GAA | AAT | GTG | ACA | 153 |
| | Asp | Pro | Asn | Pro | Gln | Glu | Val | Glu | Leu | Glu | Asn | Val | Thr | |
| | | 40 | | | | | 45 | | | | | 50 | | |
| 5 | GAA | AAT | TTT | AAC | ATG | TGG | AAA | AAT | AAC | ATG | GTA | GAA | CAG | 192 |
| | Glu | Asn | Phe | Asn | Met | Trp | Lys | Asn | Asn | Met | Val | Glu | Gln | |
| | | | | 55 | | | | | 60 | | | | | |
| | ATG | CAT | GGG | GAT | ATA | ATT | AGT | TTA | TGG | GAT | CAA | AGC | CTA | 231 |
| | Met | His | Gly | Asp | Ile | Ile | Ser | Leu | Trp | Asp | Gln | Ser | Leu | |
| | | 65 | | | | 70 | | | | | 75 | | | |
| 10 | AAG | CCA | TGT | GTA | AAA | TTA | ACC | CCA | CTC | TGT | GTT | ACG | TTA | 270 |
| | Lys | Pro | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Val | Thr | Leu | |
| | | | 80 | | | | | 85 | | | | | 90 | |
| | AAT | TGC | ACT | GAC | CCA | AAT | GTT | ACT | AAT | AGC | GAG | AGA | ACG | 309 |
| | Asn | Cys | Thr | Asp | Pro | Asn | Val | Thr | Asn | Ser | Glu | Arg | Thr | |
| 15 | | | | | 95 | | | | | 100 | | | | |
| | ATA | GAG | GGG | GGA | GAA | ATA | AAA | AAT | TGC | TCT | TTC | AAT | ATC | 348 |
| | Ile | Glu | Gly | Gly | Glu | Ile | Lys | Asn | Cys | Ser | Phe | Asn | Ile | |
| | | 105 | | | | | 110 | | | | | 115 | | |
| 20 | ACC | ACA | AAC | ATA | AGA | GAT | AGG | TTT | CAG | AAA | GAA | TAT | GCA | 387 |
| | Thr | Thr | Asn | Ile | Arg | Asp | Arg | Phe | Gln | Lys | Glu | Tyr | Ala | |
| | | | | 120 | | | | | 125 | | | | | |
| | CTT | TTT | TAT | AAA | CTT | GAT | GTA | ATA | CCA | TTA | GGT | AAT | GAT | 426 |
| | Leu | Phe | Tyr | Lys | Leu | Asp | Val | Ile | Pro | Leu | Gly | Asn | Asp | |
| | | 130 | | | | 135 | | | | | 140 | | | |
| 25 | AAT | ACT | AGC | TAT | AGG | TTG | ATA | AGT | TGT | AAC | ACC | TCA | GTC | 465 |
| | Asn | Thr | Ser | Tyr | Arg | Leu | Ile | Ser | Cys | Asn | Thr | Ser | Val | |
| | | | 145 | | | | 150 | | | | | 155 | | |
| | ATT | ACA | CAG | GCC | TGT | CCA | AAG | GTA | TCC | TTT | GAG | CCA | ATT | 504 |
| | Ile | Thr | Gln | Ala | Cys | Pro | Lys | Val | Ser | Phe | Glu | Pro | Ile | |
| 30 | | | | | 160 | | | | | 165 | | | | |
| | CCC | ATA | CAT | TAT | TGT | GCC | CCG | GCT | GGT | TTT | GCG | ATT | CTA | 543 |
| | Pro | Ile | His | Tyr | Cys | Ala | Pro | Ala | Gly | Phe | Ala | Ile | Leu | |
| | | 170 | | | | | 175 | | | | | 180 | | |
| 35 | AAG | TGT | AAA | GAT | AAG | AAG | TTC | AAT | GGA | ACA | GGA | CCA | TGT | 582 |
| | Lys | Cys | Lys | Asp | Lys | Lys | Phe | Asn | Gly | Thr | Gly | Pro | Cys | |
| | | | 185 | | | | | 190 | | | | | | |
| | ACA | AAT | GTC | AGC | ACA | GTA | CAA | TGT | ACA | CAT | GGA | ATT | AAG | 621 |
| | Thr | Asn | Val | Ser | Thr | Val | Gln | Cys | Thr | His | Gly | Ile | Lys | |
| | | 195 | | | | 200 | | | | | 205 | | | |
| 40 | CCA | GTA | GTA | TCA | ACT | CAA | CTG | TTG | TTA | AAT | GGC | AGT | CTA | 660 |
| | Pro | Val | Val | Ser | Thr | Gln | Leu | Leu | Asn | Gly | Ser | Leu | | |
| | | | 210 | | | | 215 | | | | | 220 | | |
| | GCA | GAA | GAA | GAC | ATA | GTA | ATT | AGA | TCC | GCC | AAT | CTC | ACA | 699 |
| | Ala | Glu | Glu | Asp | Ile | Val | Ile | Arg | Ser | Ala | Asn | Leu | Thr | |
| 45 | | | | | 225 | | | | | 230 | | | | |
| | GAC | AAT | GCT | AAA | AAC | ATA | ATA | GTA | CAG | CTG | AAT | GAA | TCT | 738 |
| | Asp | Asn | Ala | Lys | Asn | Ile | Ile | Val | Gln | Leu | Asn | Glu | Ser | |
| | | 235 | | | | 240 | | | | | | 245 | | |
| 50 | GTA | ACA | ATG | AAT | TGT | ACA | AGA | CCC | AAC | AAC | AAT | ACA | ATG | 777 |
| | Val | Thr | Met | Asn | Cys | Thr | Arg | Pro | Asn | Asn | Asn | Thr | Met | |
| | | | | 250 | | | | 255 | | | | | | |
| | AAA | AGT | ATA | CAT | ATA | GGA | CCA | GCG | AGA | GCA | TTT | TAT | GCA | 816 |
| | Lys | Ser | Ile | His | Ile | Gly | Pro | Gly | Arg | Ala | Phe | Tyr | Ala | |
| | | 260 | | | | 265 | | | | | 270 | | | |
| 55 | ACA | GGA | AAC | ATA | ATA | GGA | GAT | ATA | AGA | CAA | GCA | CAT | TGT | 855 |
| | Thr | Gly | Asn | Ile | Ile | Gly | Asp | Ile | Arg | Gln | Ala | His | Cys | |
| | | | 275 | | | | 280 | | | | | 285 | | |
| | AAC | ATT | AGT | GGA | ACA | AAA | TGG | AAT | GAC | ACT | TTG | AAA | AAG | 894 |
| | Asn | Ile | Ser | Gly | Thr | Lys | Trp | Asn | Asp | Thr | Leu | Lys | Lys | |
| | | | | 290 | | | | 295 | | | | | | |
| 60 | ATA | GCT | ATA | AAA | TTA | AGA | GAA | CAA | TTT | AAT | AAG | ACA | ATA | 933 |
| | Ile | Ala | Ile | Lys | Leu | Arg | Glu | Gln | Phe | Asn | Lys | Thr | Ile | |
| | | 300 | | | | 305 | | | | | | 310 | | |

| | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|------|
| | GTC | TTT | AAT | CAA | TCC | TCA | GGA | GGG | GAC | CCA | GAA | ATT | GCA | 972 |
| | Val | Phe | Asn | Gln | Ser | Ser | Gly | Gly | Asp | Pro | Glu | Ile | Ala | |
| | | | | 315 | | | | | 320 | | | | | |
| 5 | ACG | CTC | AGT | TTT | AAT | TGT | GGA | GGG | GAA | TTT | TTC | TAC | TGT | 1011 |
| | Thr | Leu | Ser | Phe | Asn | Cys | Gly | Gly | Glu | Phe | Phe | Tyr | Cys | |
| | 325 | | | | | 330 | | | | | 335 | | | |
| | AAT | TCA | ACA | CAA | CTG | TTT | AAT | AGT | ACT | TGG | AAT | AGT | ACT | 1050 |
| | Asn | Ser | Thr | Gln | Leu | Phe | Asn | Ser | Thr | Trp | Asn | Ser | Thr | |
| | | | | 340 | | | | | 345 | | | | 350 | |
| 10 | GGG | TCA | AAT | AAC | ACT | AAA | GGA | AAT | GAC | ACA | ATC | ACA | CTC | 1089 |
| | Gly | Ser | Asn | Asn | Thr | Lys | Gly | Asn | Asp | Thr | Ile | Thr | Leu | |
| | | | | | | 355 | | | | 360 | | | | |
| | CCA | TGC | AGA | ATA | AGA | CAA | ATT | ATA | AAC | ATG | TGG | CAG | AAA | 1128 |
| | Pro | Cys | Arg | Ile | Arg | Gln | Ile | Ile | Asn | Met | Trp | Gln | Lys | |
| 15 | | | | | | | | | | | | | | |
| | 365 | | | | | | 370 | | | | | 375 | | |
| | ATA | GGA | AAA | GCA | ATG | TAT | GCC | CCT | CCC | ATC | AAA | GGG | CAA | 1167 |
| | Ile | Gly | Lys | Ala | Met | Tyr | Ala | Pro | Pro | Ile | Lys | Gly | Gln | |
| | | | | | | 380 | | | | 385 | | | | |
| 20 | ATT | AGA | TGT | TCA | TCA | AAT | ATT | ACA | GGG | CTA | ATA | TTA | ACA | 1206 |
| | Ile | Arg | Cys | Ser | Ser | Asn | Ile | Thr | Gly | Leu | Ile | Leu | Thr | |
| | 390 | | | | | 395 | | | | | 400 | | | |
| | AGA | GAT | GGT | GGT | AAC | AAC | AAC | ATG | AGC | AAG | ACC | ACC | GAG | 1245 |
| | Arg | Asp | Gly | Gly | Asn | Asn | Asn | Met | Ser | Lys | Thr | Thr | Glu | |
| | | | | | | | | 410 | | | | | 415 | |
| 25 | ACC | TTC | AGA | CCT | GGA | GGA | GGA | GAT | ATG | AGG | GAC | AAT | TGG | 1284 |
| | Thr | Phe | Arg | Pro | Gly | Gly | Gly | Asp | Met | Arg | Asp | Asn | Trp | |
| | | | | | | 420 | | | | 425 | | | | |
| | AGA | AGT | GAA | TTA | TAT | AAA | TAT | AAA | GTA | GTA | AAA | ATT | GAA | 1323 |
| | Arg | Ser | Glu | Leu | Tyr | Lys | Tyr | Lys | Val | Val | Lys | Ile | Glu | |
| 30 | | | | | | | | 435 | | | | 440 | | |
| | CCA | TTA | GGA | GTA | GCA | CCC | ACC | AGG | GCA | AAG | AGA | AGA | GTG | 1362 |
| | Pro | Leu | Gly | Val | Ala | Pro | Thr | Arg | Ala | Lys | Arg | Arg | Val | |
| | | | | | | 445 | | | | 450 | | | | |
| 35 | GTG | CAG | AGA | GAA | AAA | AGA | GCA | GTG | GGA | ATA | GGA | GCT | GTG | 1401 |
| | Val | Gln | Arg | Glu | Lys | Arg | Ala | Val | Gly | Ile | Gly | Ala | Val | |
| | 455 | | | | | 460 | | | | | 465 | | | |
| | TTC | CTT | GGG | TTC | TTG | GGA | TAA | AGC | TTC | TAG | A | | | 1435 |
| | Phe | Leu | Gly | Phe | Leu | Gly | Ala | Xaa | Ser | Phe | Xaa | | | |
| | | | | 470 | | | | 475 | | | 478 | | | |

40

In addition to the listing in Table 1, Figure 3 shows the alignment of the amino acid sequences of the clones of each of the seven isolates. Corresponding residues from various clones are in boxes. In the figure, the amino acid sequences are aligned against MN-rgp120 (SEQ. ID. NO. 29).

In one embodiment, a gp120 polypeptide of this invention has the same amino acid sequence as the sequence of one of the breakthrough isolates. In another embodiment, the amino acid sequence is truncated, as described in detail hereinafter. In another embodiment, a gp120 polypeptide sequence of this invention contains a substitution, insertion, or

deletion (alteration) of one or more amino acids in the sequence of a breakthrough isolate. Usually, with the exception of amino acids that are not present in a truncated amino acid sequence and eliminate an epitope, a gp120 polypeptide of this invention will include alterations in the amino acid sequence of a breakthrough isolate that do not alter the polypeptide's ability to induce the same neutralizing antibodies as the amino acid sequence of the isolate.

In general, substitutions in the amino acid sequence of a gp120 polypeptide of this invention are conservative substitutions, particularly for amino acid residues in the V2, V3, and C4 domains of gp120, which domains contain neutralizing epitopes. However, non-conservative substitutions, particularly in domains that do not contain neutralizing epitopes are contemplated.

Conservative substitutions replace an amino acid with an amino acid of similar size and character. For example, a hydrophobic residue or hydrophilic residue is replaced with another hydrophobic residue or hydrophilic residue, respectively. Amino acids can be divided into the following groups: positively charged residues (K, R and H); negatively charged residues (D and E); amides (N and Q); aromatics (F, Y, and W); hydrophobics (P, G, A, V, L, I, and M); and uncharged residues (S and T). Usually, residues within a group are replaced with another member of the group.

In one embodiment, critical amino acid residues in the V2, V3, and C4 domains of gp120 are identical to the corresponding residues in a breakthrough isolate sequence. Critical amino acid residues in the V2, V3, and C4 domains of gp120 are described in the experimental section. In another embodiment, all amino

acid residues in the V2, V3, and C4 domains of gp120 are identical to corresponding residues in a breakthrough isolate sequence.

5 Oligonucleotide Encoding gp120 from Breakthrough Isolates

The present invention also provides novel oligonucleotides encoding gp120 from the breakthrough isolates which can be used to express gp120. An
10 oligonucleotide of this invention encodes a polypeptide of this invention. The oligonucleotide can be DNA or RNA, usually DNA. Although numerous nucleotide sequences can encode the same amino acid sequence due to the degeneracy of the genetic code, conveniently,
15 the oligonucleotides of this invention include a nucleotide sequence of a breakthrough isolate as illustrated in Table 1 (Sequence ID Nos. 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, and 28). Usually, an oligonucleotide of this invention is less than about
20 5 kilobases (kb), preferably less than about 3 kb.

To express the encoded amino acid sequence, the oligonucleotide can be inserted into a transcription unit. The transcription unit can be inserted into a plasmid for production of cell lines, inserted into a
25 virus (e.g.; vaccinia) or can be used directly as a DNA vaccine. Suitable transcription units for production of vaccine proteins are well known. A preferred expression vector, designated psvI6B5, is illustrated in Sequence ID No. 32. The vector includes an HSV-1
30 gD1 signal sequence joined to a linker sequence. The gp120 nucleotide sequence to be expressed starts with the Kpn I site of the gene. Since all gp120 or gp160 sequences contain this site, any gp120 nucleotide sequence can be analogously inserted into the vector
35 and expressed. The vector ends with a polyA tail from SV40.

In addition to being useful to express a polypeptide sequence of this invention, the oligonucleotides of this invention can also be used in diagnostics to detect HIV isolates. For example, the oligonucleotide or a portion thereof encoding a neutralizing epitope can be used in branched chain DNA diagnostics or as a probe in in situ hybridization studies.

10 Vaccine preparation

A gp120 polypeptide of this invention from a selected breakthrough isolate(s) in a suitable carrier is used to make a subunit vaccine. The polypeptide can be used alone, but is generally administered in a multivalent subunit vaccine that includes gp120 MN. In addition to one or more gp120 polypeptides of this invention, the vaccine generally includes the MN polypeptide (hereinafter, MN-rgp120). The vaccine usually includes about 3 to about 5 different gp120 polypeptides, but 30 or more different gp120 polypeptides can be used.

Preparation of gp120 polypeptides for use in a vaccine is well known and is described hereinafter. With the exception of the use of the selected HIV isolate, the gp120 subunit vaccine prepared in the method does not differ from gp120 subunit vaccines of the prior art.

As with prior art gp120 subunit vaccines, gp120 at the desired degree of purity and at a sufficient concentration to induce antibody formation is mixed with a physiologically acceptable carrier. A physiologically acceptable carrier is nontoxic to a recipient at the dosage and concentration employed in the vaccine. Generally, the vaccine is formulated for injection, usually intramuscular or subcutaneous injection. Suitable carriers for injection include

sterile water, but preferably are physiologic salt solutions, such as normal saline or buffered salt solutions such as phosphate-buffered saline or ringer's lactate. The vaccine generally contains an adjuvant.

- 5 Useful adjuvants include QS21 (Quillaja saponaria, commercially available from Cambridge Biotech, Worcester, MA), which stimulates cytotoxic T-cells, and alum (aluminum hydroxide adjuvant). Formulations with different adjuvants which enhance cellular or local
- 10 immunity can also be used. In particular, immunopotentiators such as cytokines can be included in the vaccine. Examples of suitable immunopotentiating cytokines include interleukins, such as interleukin-2 (IL-2) and interleukin-12 (IL-12), and tumor necrosis
- 15 factor-alpha (TNF- α).

- Additional excipients that can be present in the vaccine include low molecular weight polypeptides (less than about 10 residues), proteins, amino acids, carbohydrates including glucose or dextrans, chelating
- 20 agents such as EDTA, and other excipients that stabilize the protein or inhibit growth of microorganisms.

- The vaccine can also contain other HIV proteins. In particular, gp41 or the extracellular portion of
- 25 gp41 or HIV-1 core proteins such as P24, P17, and P55 can be present in the vaccine. Although the amino acid sequence of gp41 is more conserved than that of gp120, gp41 contains neutralizing epitopes. Preferably, any gp41 present in the vaccine is from an HIV isolate
- 30 present in the vaccine. gp160 from an isolate used in the vaccine can replace gp120 in the vaccine or be used together with gp120 from the isolate. Alternatively, gp160 from a different isolate than those in the vaccine can additionally be present in the vaccine.

- 35 Vaccines according to the invention can also contain one or more soluble gp120 polypeptide

sequences, or fragments thereof, in combination with an engineered virus specifically designed to express proteins that induce a cytotoxic T-cell response. Suitable engineered viruses are derived from, for example, Canary Pox virus, vaccinia viruses, attenuated human herpes viruses (such as, e.g., herpes simplex viruses), and Varicella Zoster. Exemplary engineered viruses are modified to express any HIV protein capable of inducing a cytotoxic T-cell response, such as those described above. Typically, immunization with the gp120/engineered virus vaccine is followed by administration of one or more doses of the gp120 polypeptide sequence(s) to boost the immune response. If desired, viruses can be engineered to express one or more gp120 polypeptide sequences of the invention, or fragments thereof, and used in vaccines with or without soluble gp120 polypeptide sequences.

Vaccine formulations generally include a total of about 300 to 600 μg of gp120, conveniently in about 1.0 ml of carrier. Preferred formulations include use of twice the weight of a gp120 polypeptide in twice as much alum. However, formulations having smaller amounts (e.g., 50 μg per dose) are also used, generally with alum or other adjuvants. The amount of gp120 for any isolate present in the vaccine will vary depending on the immunogenicity of the gp120. For example, gp120 from some strains of HIV may be less immunogenic than gp120 from the MN strain (Sequence ID No. 29). If two strains having different immunogenicity are used in combination, empirical titration of the amount of each virus would be performed to determine the percent of the gp120 of each strain in the vaccine. For isolates having similar immunogenicity, approximately equal amounts of each isolate's gp120 would be present in the vaccine. For example, in a preferred embodiment, the vaccine includes gp120 from the MN and a strain of this

invention at concentrations of about 300 μ g per strain in about 1.0 ml of carrier. When the vaccine includes gp120 from about 30 isolates, about 10 to about 50 μ g can be used. Methods of determining the relative amount of an immunogenic protein in multivalent vaccines are well known and have been used, for example, to determine relative proportions of various isolates in multivalent polio vaccines.

The vaccines of this invention are administered in the same manner as prior art HIV gp120 subunit vaccines. In particular, the vaccines are generally administered at 0, 1, and at 6, 8 or 12 months, depending on the protocol. A preferred protocol includes administration at 0, 1, 6, and 12 months. Following the immunization procedure, annual or bi-annual boosts can be administered. However, during the immunization process and thereafter, neutralizing antibody levels can be assayed and the protocol adjusted accordingly.

The vaccine is administered to uninfected individuals. In addition, the vaccine can be administered to seropositive individuals to augment immune response to the virus, as with prior art HIV vaccines. It is also contemplated that DNA encoding the strains of gp120 for the vaccine can be administered in a suitable vehicle for expression in the host. In this way, gp120 can be produced in the infected host, eliminating the need for repeated immunizations. Preparation of gp120 expression vehicles is described hereinafter.

Although the gp120 isolates described herein can be used as a vaccine as described above, the amino acid sequences can also be used alone or in combinations in the same type of formulation for use as an immunogen, to induce antibodies that recognize the isolate(s) present in the immunogen. Immunogens are formulated in

the same manner as vaccines and can include the same excipients, etc. Antibodies induced by the immunogens can be used in a diagnostic to detect the HIV strain in the immunogen or to affinity purify the strain.

5

gp120 Polypeptide Sequences and Chemokine Receptors

While CD4 is the primary cellular receptor for HIV-1, it is not sufficient for entry of HIV-1 into cells. Co-receptors required in conjunction with CD4 have been identified. These co-receptors are members of the chemokine receptor family of seven-transmembrane G-protein coupled receptors. The chemokine superfamily is subdivided into two groups based on the amino terminal cysteine spacing. The CXC chemokines are primarily involved in neutrophil-mediated inflammation, and the CC chemokines tend to be involved in chronic inflammation. At least five CC chemokine receptors, designated CC-CKR1-5 (also known in the art as CCR1-5), and at least four CXC chemokine receptors, designated CXC-CKR1-4 (also known as CXCR-1-4), have been identified.

CXC-CKR-4 (CXCR-4), which has also been called the alpha-chemokine receptor fusin, serves as an entry cofactor for T-cell-tropic HIV-1 strains. CC-CKR-5 (CC-R5), which has been called beta-chemokine receptor, together with its related family members, such as CC-CKR-2b and CC-CKR3, serve as entry cofactors for macrophage-tropic HIV-1 strains. T-cell-tropic strains can infect primary T-cells and T-cell lines, but not macrophages, whereas macrophage-tropic strains can infect macrophages and primary T-cells, but not T-cell lines. T-cell- and macrophage-tropic strains are discussed more fully in Deng et. al., Nature 381:661-666 (1996), which is hereby incorporated by reference in its entirety. Examples of T-cell-tropic strains include laboratory isolates, such as IIIB and MN.

Macrophage-tropic strains include primary isolates, including but not limited to A244, GNE6, GNE8, and breakthrough viruses from vaccinees immunized with gp120-based vaccines. Dual-tropic strains can use both types of co-receptors, entering cells via CXCR-4 or via one or more CC-CKR family members, preferably CC-CKR-5, CC-CKR-2b, or CC-CKR-3. While the present invention is not intended to be bound or limited by any one theory, the entry of T-cell tropic and macrophage-tropic HIV-1 strains is believed to provide a unifying explanation of the differences in cell tropism between viral strains, the resistance to HIV-1 infection by many CD4-transfected nonprimate cells, and the HIV-1-infection resistance of a portion of the human population.

Accordingly, in one embodiment is a vaccine containing (1) a first gp120 polypeptide sequence, or fragment thereof, from a macrophage-tropic HIV-1 strain and/or a second gp120 polypeptide sequence, or fragment thereof, from a T-cell tropic strain, in combination with (2) a breakthrough isolate HIV gp120 polypeptide sequence, or fragment thereof, from a vaccinee vaccinated with the first and/or second HIV gp120 polypeptide sequence. Preferably, the vaccine includes at least two gp120 polypeptide sequences that bind to different chemokine receptors. In one embodiment, the vaccine includes first and second gp120 polypeptide sequences that bind to different chemokine receptors. In addition, the breakthrough isolate gp120 polypeptide sequence can bind to a different chemokine receptor than the chemokine receptor(s) bound by either or both of the first and second gp120 polypeptide sequence(s).

A preferred T-cell tropic strain is a laboratory isolate, most preferably MN. Preferred macrophage-tropic viruses for use in the invention are GNE6 and GNE8, which are representative of the breakthrough

viruses disclosed herein and differ from MN in that their gp120s induce the formation of antibodies that recognize the gp120 sequences (e.g., the V3 domain) involved in binding to CC chemokine receptors, such as CXC-CKR-5.

In one embodiment, HIV infection is prevented by administering one or more chemokine receptor-binding gp120 polypeptide sequences, or fragment(s) thereof containing appropriate chemokine receptor-binding domains, in a vaccine, such as those described above. Preferably, the vaccine also includes one or more CD4-binding gp120 polypeptide sequences or appropriate fragments thereof. Such vaccines induce anti-HIV antibodies that inhibit viral gp120-chemokine receptor or -CD4 binding. In addition, such gp120 polypeptides can directly inhibit HIV infection by binding to one or more co-receptors for HIV infection, such as CD4 or a chemokine receptor, thus providing a prophylactic or therapeutic effect in treating HIV infection. Preferably, gp120 polypeptide sequences useful in this regard contain the T-cell binding (TCB) domain.

Various uses of chemokine receptor-binding gp120 polypeptides are discussed below with regard to the CC chemokine receptor family. However, those skilled in the art recognize that this discussion applies equally to CXC chemokine receptors that act as cofactors in HIV infection.

The gp120 polypeptides can be used as a composition containing one or more gp120 polypeptides, as described for use as a vaccine or immunogen. The composition can be administered, prophylactically or therapeutically, to a patient at risk of infection or in need of such treatment using the dosages and routes and means of administration described herein. However, chronic administration may be preferred and dosages can be adjusted accordingly. It is noted that in vivo

administration can also induce antibodies that bind viral gp120, further inhibiting virus binding to CC-CKR.

5 The gp120 polypeptides can also be used in screening assays to identify antagonists of CC-CKR. For example, candidate antagonists can be screened for inhibition of binding of gp120 to a CC-CKR CC-CKR receptor that is isolated and attached to a surface (e.g., plastic dish) or recombinantly or naturally
10 expressed on the surface of a cell. Antagonists can either bind gp120 or bind receptor. Preferred candidate antagonists include gp120 compounds, small gp120 peptides (5 to 20 amino acids in length, preferably 7 to 10 amino acids in length) or
15 peptidomimetics of gp120 that bind receptor, monoclonal antibodies that bind gp120, and small organic molecules that bind either gp120 or receptor.

The antibodies induced by the gp120 polypeptides can also be used to induce anti-idiotypic antibodies
20 that bind CC chemokines. These anti-idiotypic antibodies can be screened for binding to an anti-gp120 polypeptide antibody and inhibiting gp120 from binding CC-CKR receptor. Such anti-idiotypic antibodies mimic gp120 by binding to CC-CKR receptor. Such antibodies,
25 preferably human antibodies, can be obtained in a number of ways, such as human antibodies from combinatorial libraries (e.g., Burton et al. Adv. Immunolo. (1994) 57:191-280). It is now possible to produce transgenic animals (e.g., mice) that are
30 capable, upon immunization, of producing a full repertoire of human antibodies in the absence of endogenous immunoglobulin production. For example, homozygous deletion of the antibody heavy-chain joining region (JH) gene in chimeric and germ-line mutant mice
35 results in complete inhibition of endogenous antibody production. Transfer of the human germ-line

immunoglobulin gene array in such germ-line mutant mice results in the production of human antibodies upon antigen challenge as described in Jakobovitis et al., *Proc. Natl. Acad. Sci. USA* 90: 2551 (1993); Jakobovits et al., *Nature* 362:255-258 (1993); Bruggermann et al., *Year in Immuno.* 7: 33 (1993).

Alternatively, phage display technology as described by McCafferty et al., *Nature* 348:552-553 (1990) can be used to produce human antibodies and antibody fragments in vitro from immunoglobulin variable (V) domain gene repertoires from unimmunized donors. According to this technique, antibody V domain genes are closed in-frame either into either a major or minor coat protein gene of a filamentous bacteriophage, such as M13 or fd, and displayed as functional antibody fragments on the surface of the phage particle. Because the filamentous particle contains a single-stranded DNA copy of the phage genome, selections based on the functional properties of the antibody also result in selection of the gene encoding the antibody exhibiting those properties. Phage display can be performed in a variety of formats as reviewed by, for example, Johnson, et al., *Current Opinion in Structural Biology* 3:564-571 (1993).

Several sources of V-gene segments can be used for phage display. Clackson et al., *Nature*, 352: 624-628 (1991) isolated a diverse array of anti-oxazolone antibodies from a small random combinatorial library of V genes derived from the spleens of immunized mice. A repertoire of V genes from unimmunized human donors (or embryonic cells) can be constructed. It has been demonstrated that antibodies to a diverse array of antigens (including self-antigens) can be isolated essentially following the techniques described by Marks et al., *J. Mol. Biol.*, 222: 581-597 (1991), or Griffith et al., *EMBO J.*, 12: 725-734 (1993).

In a natural immune response, antibody genes accumulate mutations at a high rate (somatic hypermutation). Some of the changes introduced confer higher affinity, and B cells displaying high-affinity surface immunoglobulin are preferentially replicated and differentiated during subsequent antigen challenge. This natural process can be mimicked by employing the technique known as "chain shuffling" (Marks et al., *Bio/Technol.* 10:779-783 [1992]). In this method, the affinity of "primary" human antibodies obtained by phage display can be improved by sequentially replacing the heavy and light chain V region genes with repertoires of naturally occurring variants (repertoires) of V domain genes obtained from unimmunized donors. This technique allows the production of antibodies and antibody fragments with affinities in the nM range. A strategy for making very large phage antibody repertoires has been described by Waterhouse et al., *Nucl. Acids Res.*, 21: 2265-2266 (1993).

Accordingly, antibodies that bind CC-CKR can be obtained by screening antibodies or fragments thereof expressed on the surface of bacteriophage in combinatorial libraries or in other systems as described above with a gp120 monoclonal antibody that inhibits gp120 binding to receptor.

In addition to screening antibodies with a gp-120 antibody, random or combinatorial peptide libraries can be screened with either a gp120 antibody or the gp120 compounds of the invention. Approaches are available for identifying peptide ligands from libraries that comprise large collections of peptides, ranging from 1 million to 1 billion difference sequences, which can be screened using monoclonal antibodies or target molecules. The power of this technology stems from the chemical diversity of the amino acids coupled with the

large number of sequences in a library. See for example, Scott et al., *Cur. Opin. Biotechnol.* 5(1):40-8 (1994); Kenan et al. *Trends Biochem. Sci.* (1994) 19(2):57-64. Accordingly, the monoclonal antibodies, preferably human monoclonals or fragments thereof, generated as discussed herein, find use in treatment by inhibiting or treating HIV infection or disease progression, as well as in screening assays to identify additional pharmaceuticals.

Production of gp120

gp120 for a vaccine can be produced by any suitable means, as with prior art HIV gp120 subunit vaccines. Recombinantly-produced or chemically synthesized gp120 is preferable to gp120 isolated directly from HIV for safety reasons. Methods for recombinant production of gp120 are described below.

Oligonucleotides encoding gp120 from breakthrough isolates and capable of expressing gp120 can be prepared by conventional means. For example, the nucleotide sequence can be synthesized. Alternatively, another HIV nucleotide sequence encoding gp120 can be used as a backbone and altered at any differing residues as by site-directed mutagenesis. Site-directed mutagenesis is described in Kunkel et al, *Proc. Natl. Acad. Sci. (USA)* 82:488-492 (1985) and Zoller et al, *Nuc. Acids Res.* 10:6487-6500 (1982) and is well known.

In a preferred embodiment, the nucleotide sequence is present in an expression construct containing DNA encoding gp120 under the transcriptional and translational control of a promoter for expression of the encoded protein. The promoter can be a eukaryotic promoter for expression in a mammalian cell. In cases where one wishes to expand the promoter or produce gp120 in a prokaryotic host, the promoter can be a

prokaryotic promoter. Usually a strong promoter is employed to provide high-level transcription and expression.

5 The expression construct can be part of a vector capable of stable extrachromosomal maintenance in an appropriate cellular host or may be integrated into host genomes. Normally, markers are provided with the expression construct which allow for selection of a host containing the construct. The marker can be on
10 the same or a different DNA molecule, desirably, the same DNA molecule.

The expression construct can be joined to a replication system recognized by the intended host cell. Various replication systems include viral
15 replication systems such as those from retroviruses, simian virus, bovine papilloma virus, or the like. In addition, the construct may be joined to an amplifiable gene, e.g. the DHFR gene, so that multiple copies of the gp120 DNA can be made. Introduction of the
20 construct into the host will vary depending on the construct and can be achieved by any convenient means. A wide variety of prokaryotic and eukaryotic hosts can be employed for expression of the proteins.

Preferably, the gp120 is expressed in mammalian
25 cells that provide the same glycosylation and disulfide bonds as in native gp120. Expression of gp120 and fragments of gp120 in mammalian cells as fusion proteins incorporating N-terminal sequences of Herpes Simplex Virus Type 1 (HSV-1) glycoprotein D (gD-1) is
30 described in Lasky, L. A. et al., 1986 (Neutralization of the AIDS retrovirus by antibodies to a recombinant envelope glycoprotein) Science 233: 209-212 and Haffar, O.K. et al., 1991 (The cytoplasmic tail of HIV-1 gp160 contains regions that associate with cellular
35 membranes.) Virol. 180:439-441, respectively. A preferred method for expressing gp120 is described in

the examples. In the examples, a heterologous signal sequence was used for convenient expression of the protein. However, the protein can also be expressed using the native signal sequence.

5 An isolated, purified gp120 polypeptide having one of the amino acid sequences illustrated in Table 1 can be produced by conventional methods. For example, the proteins can be chemically synthesized. In a preferred embodiment, the proteins are expressed in mammalian
10 cells using an expression construct of this invention. The expressed proteins can be purified by conventional means. A preferred purification procedure is described in the examples.

15 gp120 Fragments

The present invention also provides gp120 fragments that are suitable for use in inducing antibodies for use in a vaccine formulation. A truncated gp120 sequence, as used herein, is a fragment
20 of gp120 that is free from a portion of the intact gp120 sequence beginning at either the amino or carboxy terminus of gp120. A truncated gp120 sequence of this invention is free from the C5 domain. The C5 domain of gp120 is a major immunogenic site of the molecule.
25 However, antibodies to the region do not neutralize virus. Therefore, elimination of this portion of gp120 from immunogens used to induce antibodies for serotyping is advantageous.

In another embodiment, the truncated gp120
30 sequence is additionally free from the carboxy terminal region through about amino acid residue 453 of the gp120 V5 domain. The portion of the V5 domain remaining in the sequence provides a convenient restriction site for preparation of expression
35 constructs. However, a truncated gp120 sequence that is free from the entire gp120 V5 domain is also

suitable for use in inducing antibodies.

In addition, portions of the amino terminus of gp120 can also be eliminated from the truncated gp120 sequence. In particular, the truncated gp120 sequence
5 can be free from the gp120 signal sequence. The truncated gp120 sequence can be free from the carboxy terminus through amino acid residue 111 of the gp120 C1 domain, eliminating most of the C1 domain but preserving a convenient restriction site. However, the
10 portion of the C1 domain through the V2 cysteine residue that forms a disulfide bond can additionally be removed, so that the truncated gp120 sequence is free from the carboxy terminus through amino acid residue 117 of the gp120 C1 domain. In a preferred embodiment,
15 the truncated gp120 sequence is free from the amino terminus of gp120 through residue 111 of the C1 domain and residue 453 through the carboxy terminus of gp120.

The truncated gp120 sequences can be produced by recombinant engineering, as described previously.
20 Conveniently, DNA encoding the truncated gp120 sequence is joined to a heterologous DNA sequence encoding a signal sequence.

It is understood that the application of the teachings of the present invention to a specific
25 problem or situation is within the capabilities of one having ordinary skill in the art in light of the teachings contained herein. Examples of the products of the present invention and representative processes for their isolation, use, and manufacture appear below,
30 but should not be construed to limit the invention. All literature citations herein are expressly incorporated by reference.

35

EXAMPLES

Materials and Methods

- Specimen collection from human volunteers. Blood was collected from MN-rgp120-immunized individuals who were infected with HIV-1 while participating in Phase I (NIH Protocol AVEG 016) and Phase II (NIH Protocol AVEG 201) HIV-1 vaccine trials sponsored by the National Institutes of Health (NIH). The demographics of the subjects in the study, and the study design have been described in McElrath; *Seminars in Cancer Biol.* 6:1-11 (1995); McElrath et al.; *Abstracts from Eighth Annual Meeting of the National Cooperative Vaccine Development Groups for AIDS. Bethesda, MD* 216 (1996). Specimens were obtained according to an informed consent protocol approved by the institutional review boards of the participating institutions. In the experimental section, the time of HIV-1 infection is specified with regard to data provided by the NIH AIDS Vaccine Evaluation Network where PCR (RNA) and/or serologic assays were used to detect HIV-1 infection.
- Sample preparation for cloning HIV-1 envelope glycoproteins. Peripheral blood mononuclear cells (PBMCs) from HIV-1 infected vaccinees were prepared from heparinized venous blood by FICOLL-HYPAQUE gradient centrifugation. Cell number and viability were determined. After separation, PBMCs were washed twice in phosphate-buffered saline and suspended at a cell density of 6×10^6 cells/ml in PCR lysis buffer (50 mM KCl, 10 mM Tris (pH 8.4), 2.5 mM MgCl₂, 0.1 mg/ml gelatin (Sigma), 0.45% NONIDET P40 detergent, 0.45% TWEEN 20 detergent (both detergents are commercially available from United States Biochemical Corp.) and 0.06 mg/ml Proteinase K (Gibco BRL) to lyse the cells. The lysate was incubated at 50-60°C for 1 hour, followed by inactivation of the Proteinase K at 95°C for 10 minutes. Lysates were shipped frozen and stored at -70°C until use.

Polymerase chain reaction (PCR) amplification.

Samples were subjected to two rounds of PCR amplification using the nested primers described below.

In the first round, 25 μ l aliquots of PBMC lysates (containing about 1 μ g genomic DNA) were mixed with an equal volume of a PCR reaction mix containing 400 μ M each dNTP, 200 μ g/ml BSA (Sigma Chemical Corporation, RIA grade) and about 100 pmoles of each primer in 50 mM KCl, 20 mM Tris (pH 8.4) and 3 mM MgCl₂. After an initial 10 minute denaturation step at 95°C, 5 units of Taq polymerase (AMPLITAQ, Perkin Elmer Cetus) were added during an 55°C soak step, and samples were overlaid with mineral oil.

The PCR profile was as follows: 2 cycles having 1 minute at 55°C, 2.5 minutes at 72°C and 1 minute at 94°C, followed by 28 cycles with 30 seconds at 55°C, 2.5 minutes at 72°C and 45 seconds at 94°C, and an extension step at 72°C for 5 minutes.

Aliquots of 10 μ l from the first-round reactions were re-amplified with appropriate nested primers in a final reaction volume of 100 μ l, using either the reagents and profile described above or the reagents and profile described in the PCR Optimizer Kit (Invitrogen.) PCR reaction products were purified using QIAQUICK-spin columns (Qiagen Inc.) The primer pair used in the first round was either 120.os.F (5'-gggaattcggatccAGAGCAGAAGACAGTGGCAATGA with homologous sequence at position 6248-6270 of HIVPV22) (SEQ. ID. NO. 34) or JM11A (5'-ctcgag-CTCCTGAAGACAGTCAGACTCATCAAG at position 6048-6074) (SEQ. ID. NO. 35) in the forward direction [Kusumi et al.; *J. Virol.* 66:875 (1992)] combined with 120.os.R (5'-ggctctagaagctttaGCCCATAGTGCTTCCTGCTGCT-CC at position 7836-7859) (SEQ. ID. NO. 36) in the reverse direction. The internal nested primers were 120.BX.F (5'-gggcggatcctcgaGGTACCTGTRTGGAAGAAGCA at position

6389-6410; R: A or G) (SEQ. ID. NO. 37) and 120.is.R
(5'-ggctctagaagcttttaTGCTCCYAAGAACCCAAGGAACA at position
7819-7841; Y: T or C) (SEQ. ID. NO. 38). Heterologous
primer sequences are shown in lower case letters.

5

Subcloning of PCR products and the expression of
recombinant envelope glycoproteins as fusion proteins.
The HIV-1 envelope glycoprotein gp120 sequences were
cloned and expressed as chimeric genes and fusion
10 proteins, where the signal sequence and 27 amino acids
from the mature N terminus of herpes simplex virus
type 1 (HSV-1) were fused to the N-terminal sequences
of the gp120 genes, corresponding to amino acid 13 of
the mature gp120 sequence. PCR products containing
15 gp120 sequences from the breakthrough specimens were
cloned into pRK5 expression plasmid as chimeric genes
using combinations of restrictions sites engineered
into the heterologous PCR primer tails and the Xho I
site engineered into the N-terminal sequence of
20 HSV-1 gD.

The resulting double-stranded DNA was sequenced
with Sequenase and the dGTP Reagent Kit (United States
Biochemical Corp.). Sequences from glycoprotein D were
provided to enhance expression and to provide a flag
25 epitope to facilitate protein analysis, as described in
Berman et al.; *J. Virol.* 7:4464-9 (1992); Nakamura et
al.; *AIDS and Human Retroviruses* 8:1875-85 (1992); and
Nakamura et al.; *J. Virol.* 67:6179-91 (1993).

Briefly, isolated DNA fragments generated by the
30 PCR reaction were ligated into a plasmid (pRK.gD-5,
pRKgDstop) designed to fuse the gp120 fragments, in
frame, to the 5' sequences of the glycoprotein D (gD)
gene of Type 1 Herpes Simplex Virus (gD-1) and the 3'
end to translational stop codons. The fragment of the
35 gD-1 gene encoded the signal sequence and 25 amino
acids of the mature form of HSV-1 protein. To allow

for expression in mammalian cells, chimeric genes fragments were cloned into the pRK5 expression plasmid (Eaton et al., *Biochemistry* 291:8343-8347 (1986)) that contained a polylinker with cloning sites and
5 translational stop codons located between a cytomegalovirus promotor and a simian virus 40 virus polyadenylation site.

The resulting plasmids were transfected into the 293s embryonic human kidney cell line (Graham et al.,
10 *J. Gen. Virol.* 36:59-77 (1977)) using a calcium phosphate technique (Graham et al., *Virology* 52:456-467 (1973)). Growth conditioned cell culture media was collected 48 hr after transfection, and the soluble proteins were detected by ELISA or by specific
15 radioimmunoprecipitation where metabolically labeled proteins from cell culture supernatants were resolved by sodium dodecyl sulfate polyacrylamide gel electrophoresis (PAGE) and visualized by autoradiography as described in Berman et al.,
20 *J. Virol.* 63:3489-3498 (1989) and Laemmli, *Nature* 227:680-685 (1970).

Serologic assays. Sera were assayed for antibodies to rgp120, antibodies to synthetic gp120 V3
25 domain peptides corresponding to sequences from the gp120 V3 domain, and antibodies able to inhibit the binding of MN-rgp120 to cell surface CD4 using serologic assays described in Berman et al.; *J. Virol.* 7:4464-9 (1992); Nakamura et al.; *AIDS and*
30 *Human Retroviruses* 8:1875-85 (1992); and Nakamura et al.; *J. Virol.* 67:6179-91 (1993). Endpoint titers of antibody binding to gp120 and V3 peptides were determined using three fold-serial dilutions of sera. The endpoint dilution titer was defined as the last
35 dilution that produced an optical density value that was two times higher than the mean of the optical

densities of 1:50 diluted, pooled, normal human sera. Antibody titers were calculated by a computer program that interpolated values between antibody dilutions. The inter-assay coefficient of variation of positive
5 control standard sera was 35%.

Binding of monoclonal antibodies to rgp120 from breakthrough viruses. An ELISA similar to that described by Moore et al.; *AIDS* 3:155-63 (1989) was
10 used to measure the binding of various monoclonal antibodies (MAbs) to rgp120s from breakthrough viruses. Briefly, Nunc-Immuno plates (Maxisorp, certified) were coated (100 μ l at 5 μ g/ml in PBS at 4°C overnight) with an affinity-purified sheep polyclonal antiserum to a
15 peptide at the C terminus of gp120 (D7324, International Enzymes, Fallbrook, CA). After washing once with PBS-0.05% TWEEN-20 detergent, the plates were blocked with PBS-1.0% BSA for 30-60 minutes at room temperature. Cell culture supernatants from 293s
20 cells, diluted to contain equivalent amounts of the gD-rgp120 fusion protein, were added and incubated for 2 hours at room temperature followed by three washes with PBS-0.05% TWEEN-20 detergent. Various MAbs were diluted in PBS-1.0% BSA and 100 μ L of the diluted MAbs
25 were added to each well and incubated for 1 hour at room temperature.

The plates were washed 3 times and incubated with 100 μ l of a horseradish peroxidase-conjugated second antibody (goat anti-mouse or anti-human IgG, Cappel)
30 for 1 hour at room temperature. After 3 washes the plates were developed and the OD₄₉₂ (optical density at 492 nm) read in a plate reader. Growth conditioned cell culture supernatants were normalized by dilution based on binding by MAb 5B6 which is specific for HSV-1
35 glycoprotein D fusion protein.

Virus neutralization assays. The ability of vaccinee sera to inhibit infection of MT4 cells by HIV-1_{MN} was measured in a cytopathicity assay where cell viability was quantitated using a calorimetric indicator dye, as described in Robertson et al.; *J. Virol. Methods* 20:195-202 (1988). Briefly, a virus stock of HIV-1_{MN} (obtained from Dr. Michael Norcross, U.S. Food and Drug Administration) was prepared as the clarified supernatant from chronically infected H9/HIV-1_{MN} cell culture. H9 cells chronically infected with HIV-MN were pelleted and resuspended in one-tenth the original volume of medium. Cell-associated virus was released by the mechanical shearing effects of rapid vortexing of the cells as described in Wrin et al.; *J. Virol.* 69:39-48 (1995).

An amount of virus sufficient to ensure complete cell lysis killing in 7 days was incubated with three-fold serial dilutions of test antisera, and then used to challenge MT4 T-lymphoid cells in 10% FCS/RPMI-1640 cell culture media. The cultures were incubated for 7 days at 37°C in 5% CO₂, and then cell viability was tested by the dye MTT, as described by Robertson et al.; *J. Virol. Methods* 20:195-202 (1988). Virus neutralization endpoints were quantitated by measurement of OD at 570-650 nm, and then the endpoint titers were calculated as the reciprocal of the antiserum dilution giving a signal that was two-fold above the control signal with unprotected (killed) cells. These titers were typically twice those calculated at 50% protection.

Results

Immunization history of infected subjects. Since 1992, 499 adults have been immunized with MN-rgp120 in Phase I trials in low or moderate risk individuals and in a Phase II clinical trial involving moderate to high

5 risk individuals. The studies described herein entail the genetic and immunologic characterization of the first seven of nine individuals who became infected with HIV-1 through high risk behavior during the course of these trials. A listing of the trials and summary of the status of the vaccinees is presented in Table 2A. A listing of the analysis of the vaccinees is presented in Table 2B.

10

TABLE 2A

Description of Vaccinees Infected with HIV-1
After Immunization with MN-rgp120

| | <u>Study No.</u> | <u>Case No.</u> | <u>*Risk Group</u> | <u>‡Antigen dose/ Adjuvant</u> |
|----|------------------|-----------------|--------------------|------------------------------------|
| 15 | 016 | C6 | M/H | 300/QS21 |
| | 016 | C8 | M/H | 600/QS21 |
| | 016 | C15 | M/H | 300/QS21 |
| | 201 | C7 | M/H | 600/Alum |
| | 201 | C11 | M/H | 600/Alum |
| 20 | 201 | C10 | M/IDU | 600/Alum |
| | 201 | C17 | M/IDU | 600/Alum |

* - M/H indicates male homosexual; M/IDU indicate male intravenous drug user.

‡ - numbers indicate dose in micrograms of MN-rgp120 injected per immunization; QS21 indicates antigen was formulated in QS21 adjuvant; Alum indicates MN-rgp120 formulated in aluminum hydroxide.

25

TABLE 2B
Description of Vaccines Infected with HIV-1
After Immunization with MN-rgp120

| 5 | Case No. | Injection Schedule | Injections before HIV-1+ | Time of HIV-1+ (months) | Interval: to HIV-1+ (months) |
|----|----------|--------------------|--------------------------|-------------------------|------------------------------|
| | | (months) | HIV-1+ | (months) | (months) |
| | C6 | 0,1,10.5 | 2 | 4.00 | 2.00 |
| | C8 | 0,1 | 2 | 4.00 | 3.00 |
| | C15 | 0,1,2 | 3 | 6.25 | 4.00 |
| 10 | C7 | 0,1,6,12 | 3 | 9.25 | 3.00 |
| | C11 | 0,1,6,12 | 4 | 19.50 | 6.75 |
| | C10 | 0,1,6,19 | 3 | 19.50 | 13.50 |
| | C17 | 0,1,6,18 | 4 | 24.75 | 6.25 |

□ - indicates interval between last immunization and detection of HIV-1 infection.

15

Three of the infections occurred in a Phase I trial (NIH Protocol AVEG 201) that compared the safety and immunogenicity of MN-rgp120 formulated in two different adjuvants (alum and QS21), and four of the infections occurred in a Phase II trial aimed at establishing the safety and immunogenicity of MN-rgp120 in various high risk groups (e.g., intravenous drug users, homosexual and bisexual males, and partners of HIV-1 infected individuals).

Of the seven infections studied (Table 3), two (C6 and C8) occurred after two injections, three (C7, C10 and C15) occurred after three injections, and two (C11 and C17) occurred after receiving the four scheduled injections. The interval between receiving the last immunization and becoming infected was 2 to 13.5 months.

TABLE 3

**Pak P st Boost MN-rgp120 Antibody Titers
in Vaccinees that Became Infected with HIV-1**

| 5 | <u>Injections</u> | <u>C6</u> | <u>C8</u> | <u>C15</u> | <u>C7</u> | <u>C11</u> | <u>C10</u> | <u>C17</u> |
|----------|--------------------------|------------------|------------------|-------------------|------------------|-------------------|-------------------|-------------------|
| | 1 | <50 | 2185 | 79 | <50 | 1890 | na | na |
| | 2 | 21539 | 10125 | na | 413 | 32696 | 7771 | 7056 |
| | 3 | # | # | 4460 | 9707 | 34728 | 11627 | 1841 3 |
| | 4 | # | # | # | # | # | # | 1134 0 |

10 # - indicates specimen not analyzed because of HIV-1 infection.
na - indicates the sample was not available for testing.
15 boldface - indicates unusually low antibody titers.

Antibody response to gp120 in vaccinated individuals. The magnitude and specificity of the antibody response to MN-rgp120 was measured by ELISA in
20 all infected individuals throughout the course of the immunization regime (Figure 1). Five of the seven

subjects exhibited normal antibody response kinetics that included a small but reproducible primary response (1:100-1:2,000) and a strong secondary (booster) response (titers ranging from 1:7,000-1:32,000), and
5 antibody responses following third and fourth injections that were similar or marginally higher than those achieved after the second immunization (Figure 1, Table 3).

The antibody response observed in C7 (Figure 1C)
10 was unusual in that no antibodies were detectable after the primary injection and a titer of only 1:350 was detected after the second injection. It thus appeared that C7 did not respond to the primary immunization, and that the antibody response obtained after the
15 second injection represented a primary immune response. Consistent with this hypothesis, the third injection elicited a titer of only 1:9,707, typical of those normally seen after two immunizations.

An atypical antibody response was also seen in
20 subject C15 (Figure 1G) who was immunized according to an accelerated immunization schedule of 0, 1, and 2 months. As expected, the antibody titer seen in this subject (1:4,460) was at the low end of what is typically achieved after two immunizations and was far
25 below normal values for three immunizations. The lack of an effective booster response after the third immunization of C15 was not surprising in view of previous studies where an accelerated 0, 1, and 2 month immunization schedule in baboons [Anderson et al.;
30 *J. Infect. Dis.* 160:960-9 ((1989))] similarly prolonged the secondary response and failed to elicit an effective tertiary booster response.

Retrospective analysis of serum and plasma from subjects C6 (Figure 1A) and C8 (Figure 1B) indicated
35 that they became infected with HIV-1 at some point between the second and third immunizations. Serologic

evidence of HIV-1 infection was evident in the gp120 antibody assays where the titers failed to decline two weeks after the second injection and instead formed an uncharacteristic high titer plateau (Figures 1A and 1B). A similar plateau in MN-rgp120 titer after the third injection, suggested that subject C7 became infected around week 36, approximately 16 weeks after receiving the third injection (Figure 1C). Subjects C10 (Figure 1E), C11 (Figure 1D), C15 (Figure 1G), and C17 (Figure 1F) developed unexpected increases in gp120 titers, typical of HIV-1 infection, after either the third or fourth immunizations. The data obtained demonstrate that immunologic priming for MN-rgp120 antibody responses is insufficient to provide universal protection from HIV-1 infection.

Antibody titers to the V3 domain. To further characterize the antibody response to gp120, antibody titers were measured to a synthetic V3 domain peptide of MN-rgp120 containing the principal neutralizing determinant (PND). Five of the seven subjects developed good V3 titers (1:400 to 1:4000) after the second immunization, however two subjects (C7 and C15) required three immunizations before developing significant tiers (Figures 1C and 1G). As had been observed previously (11), the peak V3 titers in some individuals (e.g. C11, C10, C17) appeared to decline with each successive immunization (Figures 1D, 1E, and 1F). After HIV-1 infection, two patterns of V3 reactivity were observed. Three subjects (C6, C7, and C10) showed large increases in titer to V3 domain peptides (Figures 1A, 1C, and 1E) whereas C8 (Figure 1B) showed a large decrease in V3 titer. At the time of analysis, the data were insufficient to

draw any conclusions regarding the changes in V3 titers in response to HIV-1 infection in subjects C11, C15 and C17.

5 The results obtained indicate that the ability to form antibodies reactive with the V3 domain at various time-points prior to HIV-1 infection is not a valid correlate of protective immunity against all strains of HIV-1.

10 CD4 Inhibition titers. Antibodies that block the binding of gp120 to CD4 represent a heterogeneous class of virus neutralizing antibodies. Some are known to bind to the C4 domain of gp120 [Nakamura et al.; *J. Virol.* 67:6179-91 (1993); Anderson et al.; *J.*
15 *Infect. Dis.* 160:960-9 ((1989)), and some are known to recognize conformation dependent discontinuous epitopes [Berman et al.; *J. Virol.* 7:4464-9 (1992); Nakamura et al.; *J. Virol.* 67:6179-91 (1993); McKeating et al.; *AIDS Research and Human Retroviruses*
20 8:451-9 (1992); Ho et al.; *J. Virol.* 65:489-93 (1991); Barbas et al.; *Proc. Natl. Acad. Sci. USA* 91:3809-13 (1994)].

One way to detect antibodies to both types of epitopes is to measure the ability of vaccinee sera to
25 prevent the binding of [¹²⁵I]-labeled gp120 to cell surface CD4 [(Nakamura et al.; *AIDS and Human Retroviruses* 8:875-85 (1992); Nakamura et al.; *J. Virol.* 67:6179-91 (1993)]. CD4 blocking titers were detected in all seven of the vaccinees prior to
30 infection (Figure 2) with peak titers that ranged from 1:10-1:300. At the last time point prior to infection, the CD4 titers in five of the seven vaccinees was low (1:30 or less). One vaccinee (C17), however, possessed a CD4 blocking titer of about 1:300 prior to infection
35 (Figure 2F). Thus, the lack of antibodies that block the binding of MN-rgp120 to CD4 cannot account for all

of the infections. Large increases in CD4 blocking titers (1:100-1:1,000) were seen in five of the seven subjects after HIV-1 infection. These included vaccinees C6, C7, C8, C10, and C11. These results demonstrate that the CD4 blocking titers elicited by MN-rgp120 were lower than those elicited by natural infection.

Virus neutralizing activity. The virus neutralizing activity of antisera from MN-rgp120-immunized subjects was measured using a colorimetric assay that measured the viability of MT-4 cells after incubation with antibody treated virus (HIV-1_{MN}). Since the actual date of infection was not known for any of the breakthrough infections, and serum samples were collected infrequently, the magnitude of the neutralizing antibody response at the time of infection is not known for any of the vaccinees.

Of the seven infections examined, the serum sample closest to the time of infection was that obtained from C7, where a neutralizing titer of 1:15 to HIV-1_{MN} was present three weeks prior to detection of HIV-1 infection (Table 4). In all other cases, however, the interval between the last injection and the time of infection was 10 to 25 weeks.

TABLE 4
Neutralization Activity of Sera from Vaccines
Infected with HIV-1

| | <u>Week</u> | <u>C6</u> | <u>C8</u> | <u>C15</u> | <u>C7</u> | <u>C11</u> | <u>C10</u> | <u>C17</u> |
|----|-------------|-----------|-----------|------------|-----------|------------|------------|------------|
| 5 | 0 | <10* | <10* | <10* | <10* | <10* | <10* | <10* |
| | 2 | <10 | <10 | <10 | - | - | - | - |
| | 4 | <10* | <10* | nd* | <10* | <10* | <10* | <10* |
| | 6 | 10 | 80 | - | <10 | 30 | 150 | 150 |
| | 8 | - | - | nd* | - | - | - | - |
| 10 | 10 | - | - | 35 | - | - | - | - |
| | 15 | - | - | - | <10 | - | - | - |
| | 16 | 150# | 250# | - | - | 30 | 10 | <10 |
| | 24 | | | 150# | <10* | 20* | <10* | <10* |
| | 26 | | | | 70 | 500 | 200 | 400 |
| 15 | 30 | | | | - | - | 40 | 100 |
| | 33 | | | | 15 | - | - | - |
| | 35 | | | | - | 100 | - | - |
| | 36 | | | | 30# | - | 10 | 40 |
| | 52 | | | | | 30* | <10 | <10 |
| 20 | 54 | | | | | 250 | - | - |
| | 57 | | | | | 100 | - | - |
| | 63 | | | | | 90 | - | - |
| | 64 | | | | | - | - | <10 |
| | 77 | | | | | 40# | - | - |
| 25 | 78 | | | | | | 500# | 10* |
| | 80 | | | | | | | 100 |
| | 84 | | | | | | | 60 |

90

150

104

150#

- 5 * - indicates immunization.
 # - indicates HIV-1 positive.
 nd - indicates not done.
 - - indicates sample not available.

10 When sera from the two early infections were
 examined (Table 4), one individual (C6) had a peak
 neutralizing titer of 1:10 ten weeks prior to detection
 of HIV-1 infection, whereas the other individual (C8)
 had a neutralizing titer of 1:80 ten weeks prior to
 detection of HIV-1 infection. Subject C15, who was
15 immunized according to an accelerated immunization
 schedule, developed a neutralizing titer of 1:35 after
 the third injection, 14 weeks prior to HIV-1 infection.
 Subject C10, who had a peak neutralizing titer of 1:200
 following the third immunization (week 24), had no
20 detectable titer at week 52, six months prior to the
 first indication of HIV-1 infection (week 78).

 Subject C11 possessed a neutralizing titer of 1:90
 at fourteen weeks prior to detection of HIV-1 and a
 peak titer of 1:500 following the third immunization.
25 Similarly vaccinee C17 had a neutralizing titer of
 1:150 fourteen weeks prior to infection and a peak
 titer of 1:400 at two weeks after the third
 immunization.

 Based on the rate of decay of the gp120 response
30 of approximately two months [Belshe et al.; JAMA
 272(6):475-80 (1994)], as well as the observation that
 neutralizing titers of 1:150 decayed to 1:10 in 10
 weeks in vaccinees C10 and C17, it appears that
 neutralizing titers in C8, C15, C11, and C17 could have
35 declined to 1:10 or less in the intervals between the
 last pre-infection serum sample and the time of HIV-1

detection.

The results of these studies demonstrated that all vaccinees developed some level of virus-neutralizing antibodies at some time prior to HIV-1 infection, and that the magnitude of the neutralizing response was probably low at the time of infection. In general, the magnitude of the virus-neutralizing response observed in the individuals that became infected with HIV-1 was comparable to that seen in non-infected vaccinees as described in Belshe et al.; *JAMA* 272(6):475-80 (1994).

Sequences of Viruses. To evaluate the similarity of the breakthrough viruses with the vaccine antigen, nucleotide sequences for gp120 from all seven breakthrough viruses were determined. Envelope glycoprotein genes were amplified from proviral DNA using the polymerase chain reaction. Sequences were obtained by direct amplification of DNA from lysates of gradient-purified lymphocytes obtained directly from patient blood without any intermediate tissue culture or amplification step.

A listing of the complete gp120 sequences (two clones per specimen) is provided in Figure 3. All seven envelope glycoproteins possessed sequences typical of subtype (clade) B viruses. The overall homology with MN-rgp120 ranged from 69-80% (Table 5).

TABLE 5
Comparison of MN-rgp120 Sequence with Sequences
from Infected Vaccinees*

| | | MN | C6.1 | C8.3 | C7.2 | C11.5 | C10.5 | C17.1 | C15.2 |
|----|-------|-----|------|------|------|-------|-------|-------|-------|
| 5 | MN | 100 | 79 | 78 | 70 | 75 | 69 | 80 | 72 |
| | C6.1 | | 100 | 78 | 70 | 81 | 75 | 90 | 79 |
| | C8.3 | | | 100 | 68 | 80 | 76 | 84 | 83 |
| | C7.2 | | | | 100 | 80 | 73 | 76 | 73 |
| | C11.5 | | | | | 100 | 75 | 70 | 80 |
| 10 | C10.5 | | | | | | 100 | 70 | 72 |
| | C17.1 | | | | | | | 100 | 87 |
| | C15.2 | | | | | | | | 100 |

* - Data indicate percent identity.

- 15 Interestingly, a high percentage (four of seven) of the breakthrough viruses differed from MN-rgp120 by 25-30% [Myers et al.; *Retroviruses and AIDS Database, Los Alamos National Laboratory* (1992 and 1995)].
- 20 Historically this degree of sequence variation is typical of inter-subtype (intra-clade) variation rather than intra-subtype variation which is expected to be in the 10-20% range [Myers et al.; *Retroviruses and AIDS Database, Los Alamos National Laboratory* (1992 and 1995)]. Of the viruses with the greatest homology to
- 25 MN-rgp120, two (C6 and C8) occurred as early infections, prior to complete immunization, and one (C17) occurred as a late infection.

- 30 **Polymorphism in the V3 Domain.** Of particular interest were polymorphisms in regions known to contain epitopes recognized by virus neutralizing antibodies. The best characterized neutralizing epitope, the principal neutralizing determinant (PND), occurs at the

tip of the V3 loop. In subtype B viruses, approximately 60% possess the MN serotype-defining signature sequence, IGPGRAF (SEQ. ID. NO. 39), based on identity with the prototypic MN strain of HIV-1 [Berman et al.; *J. Virol.* 7:4464-9 (1992); Myers et al.; *Retroviruses and AIDS Database*, Los Alamos National Laboratory (1992 and 1995); La Rosa et al.; *Science* 249:932-5 (1990)].

Three of the viruses (C6, C8, and C17) possessed the MN serotype signature sequence (Figure 3). In contrast, four viruses possessed sequences with radical amino acid substitutions in the PND [IGPGRAW (C7), LGPGSTF (C11), IGPGRVL (C10), and IGPGSAF (C15)] (SEQ. ID. NOS. 40-43, respectively), and therefore were classified as "non-MN like" viruses. Of note, each of the four "non-MN-like" sequences were rare (Table 6) and were not typical of the most common "non-MN" variants of subtype B viruses [Myers et al.; *Retroviruses and AIDS Database*, Los Alamos National Laboratory (1992 and 1995)].

TABLE 6
Frequency of Polymorphisms at the Principal
Neutralizing Determinant in HIV-1 Infected
Individuals Immunized with MN-rgp120*

| 5 | V3 Sequence | | Observed | Dataset Frequency | | | |
|----|-----------------|----------|------------------|-------------------|----------------|----------------|----------------|
| | Sequence | n | | GNE | LANL | LANL.1 | LaRosa |
| | <u>Sequence</u> | <u>n</u> | <u>Frequency</u> | <u>(n=52)</u> | <u>(n=519)</u> | <u>(n=160)</u> | <u>(n=245)</u> |
| | GPGRAPH | 3 | 0.42 | 0.67 | 0.57 | 0.66 | 0.60 |
| | GPGRW | 1 | 0.14 | 0.03 | 0.013 | 0.06 | 0.010 |
| 10 | GPGRVL | 1 | 0.14 | <0.02 | 0.004 | <0.006 | <0.008 |
| | GPGSTF** | 1 | 0.14 | <0.02 | <0.002 | <0.006 | <0.004 |
| | GPGSF | 1 | 0.14 | 0.02 | 0.011 | <0.006 | <0.004 |

15 * - Data set GNE refers to a collection of
52 independent isolates collected in 1992;
dataset LANL refers to a collection of
20 519 sequences reported by Myers et al.,
Retroviruses and AIDS Database, Los Alamos
National Laboratory 1992 and 1995; LANL.1 refers
to a collection of 160 epidemiologically unlinked
individuals provided by B. Korber (personal
communication); dataset La Rosa refers to sequence
25 data reported by La Rosa et al., *Science* 249:932-5
(1990).
** - Sequences were not present in the data sets
examined.

30 The prevalence of viruses with PND sequences
matching the breakthrough viruses ranged from a high of
1.3% (C7) to a low of 0.2% (C11) in a listing of 519
subtype B sequences compiled by the Los Alamos National
Laboratory [Myers et al.; *Retroviruses and AIDS*
35 *Database, Los Alamos National Laboratory* (1992 and
1995)]. Similarly low frequencies were observed in

three other independently derived data sets (Table 6). The occurrence of these sequences did not differ significantly between data sets collected prior to 1985 [La Rosa et al.; *Science* 249:932-5 (1990)] and data
5 collected 1992, or from a set of 160 epidemiologically unlinked individuals (B. Korber, personal communication). All four sets of data agreed that the prevalence of viruses with MN-like PND sequences was in the range of 60%. Based on this data, four of the
10 seven breakthrough infections were determined to be caused by viruses that fell outside of the spectrum of viruses that the vaccine was expected to prevent.

Other features of breakthrough virus V3 domains.

15 Like MN-rgp120, the V3 domains of all of the breakthrough viruses were 36 amino acids in length. However, all seven viruses differed from MN-rgp120 with respect to the number of glycosylation sites and with respect to the syncytium-inducing (SI) signature
20 sequence.

The sequence of MN-rgp120 is somewhat unusual [Myers et al.; *Retroviruses and AIDS Database, Los Alamos National Laboratory* (1992 and 1995)] in that it
25 lacks an N-linked glycosylation site at position 306 in the V3 domain. The lack of this glycosylation site does not appear to be antigenically significant since antisera to MN-rgp120 are known to neutralize a variety of viruses (e.g. SF-2, DU6587-5, DU4489-5, CC) that possess a glycosylation site at this position
30 [Berman et al.; *J. Virol.* 7:4464-9 (1992)]

In addition, the V3 domain of MN-rgp120 possessed sequence polymorphisms (R at position 311, K at position 324, K at position 328) typical of syncytium
35 inducing viruses [Fouchier et al.; *J. Virol.* 66:3183-87 (1992)], whereas all seven breakthrough viruses possessed sequences associated with non-syncytium-

inducing viruses. Syncytium-inducing viruses have been associated with rapid disease progression [Tersmette et al.; *J. Virol.* 62:2026-32 (1988)] and T cell tropism [O'Brien et al.; *Nature (London)* 348:69-73 (1990);
5 Shioda et al.; *Nature (London)* 349:167-9 (1991)]. To date viruses with these properties have not been recovered from any of the MN-rgp120 immunized volunteers.

10 **Polymorphism in the V1, V2 and C4 domains.**

Previous investigations have identified additional neutralizing epitopes in the V1, V2 and C4 domains of gp120 [Nakamura et al.; *J. Virol.* 67:6179-91 (1993);
McKeating et al.; *AIDS Research and Human Retroviruses*
15 8:451-9 (1992); Ho et al.; *J. Virol.* 65:489-93 (1991);
Barbas et al.; *Proc. Natl. Acad. Sci. USA* 91:3809-13 (1994);
McKeating et al.; *J. Virol.* 67:4932-44 (1993);
Moore et al.; *J. Virol.* 67:6136-6151 (1993);
Davis et al.; *J. Gen. Virol.* 74:2609-17 (1993)].

20 The best characterized of these neutralizing epitopes is in the C4 domain which has attracted special attention because antibodies binding to this area are known to block the binding of gp120 to CD4 [Moore et al.; *AIDS* 3:155-63 (1989); McKeating et al.;
25 *AIDS Research and Human Retroviruses* 8:451-9 (1992)]. Because the epitope is located in a conserved (C) domain, naturally-occurring polymorphism in this region is far more limited than in other neutralizing epitopes. Nakamura et al.; *J. Virol.* 67:6179-91 (1993)
30 reported that the binding of a number of neutralizing MAbs was dependent on K at position 429.

Comparison of the sequence of MN-rgp120 with other strains of HIV-1 showed that a common polymorphism, involving the substitution of E for K, occurs at this
35 position. Indeed, substrains of the same virus isolate often show polymorphism at this position. The HXB2

substrain of HIV-1_{LAI} contains K at position 429, whereas the BH10, IIIB, and LAV substrains of the HIV-1_{LAI} contain E at this position [Nakamura et al.; *J. Virol.* 67:6179-91 (1993)]. Similarly, the 1984 isolate of HIV-1_{MN} exhibited E at this position, while the 1990 isolate of HIV-1_{MN}, used to produce MN-rgp120, possessed K at this position.

When the sequences of the infected vaccine recipients were examined (Figure 3), the virus from subject C17, like MN-rgp120 contained K at position 429, whereas the six other viruses that differed from the vaccine immunogen possessed E at this position. These results demonstrated that six of the seven breakthrough viruses differed from the vaccine immunogen at the CD4-blocking, neutralizing epitope in the C4 domain of gp120.

Studies with monoclonal antibodies have defined neutralizing epitopes in the V1 and V2 domains of gp120 [McKeating et al.; *J. Virol.* 67:4932-44 (1993); Moore et al.; *J. Virol.* 67:6136-6151 (1993); Davis et al.; *J. Gen. Virol.* 74:2609-17 (1993)]. Like the polymorphisms that occur in the C4 domain, the V2 domains exhibit several common polymorphisms that affect the binding of virus neutralizing antibodies. One such polymorphism occurs at position 171 which is critically important for the binding of murine MAb 1025, whereas residue 187 is important for the binding of MAb several MABs represented by 1088.

When the V2 domain sequences were examined (Figure 3), all of the infected-vaccinee viruses differed from MN-rgp120 in that R replaced G at position 171 and I or V replaced E at position 187. Antibodies recognizing these adjacent sites in the V2 domain of MN-rgp120 would not be expected to neutralize viruses with radical amino acid substitutions at these position. Thus, all seven

breakthrough viruses differed from MN-rgp120 at a neutralizing epitope in the V2 domain of gp120.

Other neutralizing epitopes have been reported in the V1 domain of gp120 [O'Brien et al.; *Nature (London)* 348:69-73 (1990); McKeating et al.; *J. Virol.* 67:4932-44 (1993)]. Although the neutralizing epitopes in the V1 domain of MN-rgp120 have not been characterized, the polymorphism seen among the breakthrough viruses in this region was interesting. Particularly striking (Figure 3) was that the length of this domain ranged from 20 amino acids (C17) to 45 amino acids (C6), and the number of N-linked glycosylation sites ranged from 2 to 6. In contrast, the V1 domain of MN-rgp120 is 31 amino acids in length and encodes three N-linked glycosylation sites.

Although examination of sequence databases suggest that variation in the V2 region is comparable to the V1 region, the V2 region of the breakthrough viruses showed less variation than expected. Specifically, the length of the V2 region ranged from 36 amino acids (C7) to 39 amino acids in length, with six of seven viruses containing three N-linked glycosylation sites in this domain. A high degree of polymorphism was found in the V4 region where sequences ranged from 26 (C10) to 33 (C15, C7) amino acids in length and contained either 4 or 5 N-linked glycosylation sites.

Antigenicity of envelope glycoproteins from breakthrough viruses. To determine the significance of sequence variation on glycoprotein antigenicity, recombinant gp120 was prepared from the viruses of all seven infected vaccinees (Figure 4). In these studies a series of MAbs was assembled and their binding to MN-rgp120 was compared to that of rgp120 from the vaccinee isolates by ELISA (Table 7).

TABLE 7
Relative Reactivity* of MAb Binding to rgp120 from
Infect d Subjects Compared with Binding to MN-rgp120

| | | <u>V3</u> | | <u>Discontinuous</u> | | <u>C8</u> | <u>V2</u> |
|----|--------------|-------------|-------------|----------------------|-------------|-------------|-------------|
| 5 | <u>gp120</u> | <u>1034</u> | <u>50.1</u> | <u>1.5E</u> | <u>1025</u> | <u>1024</u> | <u>1088</u> |
| | MN | 1.0 | 1.00 | 1.00 | 1.00 | 1.00 | 1.00 |
| | C6.1 | 0.37 | 0.37 | 0.17 | 0.00 | 0.00 | 0.00 |
| | C6.5 | 0.33 | 0.33 | 0.75 | 0.00 | 0.00 | 0.00 |
| | C8.3 | 0.11 | 0.37 | 0.38 | 0.00 | 0.00 | 0.00 |
| 10 | C8.6 | 0.14 | 0.34 | 0.29 | 0.00 | 0.00 | 0.00 |
| | C7.2 | 0.47 | 0.60 | 0.71 | 0.00 | 0.00 | 0.00 |
| | C11.5 | 0.00 | 0.00 | 0.17 | 0.00 | 0.00 | 0.00 |
| | C11.7 | 0.00 | 0.00 | 0.17 | 0.00 | 0.00 | 0.00 |
| | C10.5 | 0.33 | 0.40 | 0.46 | 0.24 | 0.03 | 0.04 |
| 15 | C10.7 | 0.42 | 0.48 | 0.50 | 0.29 | 0.07 | 0.09 |
| | C17.1 | 0.33 | 0.52 | 0.33 | 0.00 | 0.30 | 0.07 |
| | C17.3 | 0.37 | 0.56 | 0.33 | 0.00 | 0.38 | 0.06 |
| | C15.2 | 0.00 | 0.47 | 0.92 | 0.00 | 0.00 | 0.00 |
| | C15.3 | 0.00 | 0.37 | 0.63 | 0.00 | 0.00 | 0.00 |

20

* - Relative reactivity values represent ratio of optical densities obtained with rgp120 from patient isolates divided by optical density obtained for MN-rgp120 at a MAb concentration of 2 micrograms per milliliter.

25

In control experiments, the binding of MAb 5B6 (which is specific for the HSV gD-1 flag epitope fused to the N terminus of all of the rgp120 protein) was used to standardize the amount of gp120 from each isolate (Figure 5A). These studies demonstrated that the assay was carried out under conditions where equivalent amount of rgp120s were captured onto wells of microtiter plates.

35

The antigenic structure of the V3 domain was examined using the 1034 MAb (isolated from mice immunized with MN-rgp120 as described in Nakamura et al.; *J. Virol.* 67:6179-91 (1993) and the 50.1 MAb (prepared from mice immunized with a synthetic V3 domain peptide as described in Rini et al.; *Proc.*

40

Natl. Acad. Sci. USA 90:6325-9 (1993). Both MAbS are known to exhibit potent virus neutralizing activity. When binding to the recombinant proteins was examined, the MAb binding to MN-rgp120 was at least 10-fold greater than to any of the breakthrough virus envelope proteins (Figure 5 B and C). Surprisingly, rgp120 from the three patient isolates (C8, C6, and C17) that possessed the MN serotype-defining sequence, IGPGRF (SEQ. ID. NO. 39), varied from one another in their MAb binding activity. Thus, the binding of MAb 1034 and MAb 50.1 to rgp120 from C17 was significantly greater than the binding to rgp120s from C6 and C8.

A distinction in the epitopes recognized by these MAbS was evident since C6-rgp120 and C8-rgp120 gave comparable binding with 50.1, whereas 1034 bound better to the C6-derived protein than the C8-derived protein. The poorest MAb reactivity was with rgp120s from C11 and C15. This result was consistent with sequence analysis demonstrating that these two viruses both possessed the radical substitution of S for R at position 18 in the V3 domain. Surprisingly, both of these MAbS exhibited better than expected binding to rgp120 from the C7 and C10 viruses. Like MN-rgp120, both proteins contained the penta-peptide, IGPGR sequence (SEQ. ID. NO. 44) in the V3 loop, but differed from MN-rgp120 in that V and L replaced A and F at positions 319 and 320 in gp120 from C10, and W replaced F at position 320 in gp120 from C7. These results indicate that R at position 318 is essential for the binding of these two MAbS, and that the epitopes recognized by 1034 and 50.1 are not completely destroyed by the hydrophobic substitutions at positions 319 and 320.

As predicted from the sequence data, there was little if any binding to the breakthrough virus rgp120s using MAbS (1088 and 1025) directed to the V2 region of

MN-rgp120. Also consistent with sequence data was the observation that MAb 1024 directed to the C4 domain of MN-rgp120 gave some reactivity with C17-rgp120 which, like MN-rgp120 contained K at position 429, but gave no reactivity with the other isolates that contained E at residue 429.

Together, these studies demonstrated that the antigenic structure of all seven breakthrough viruses differed from the vaccine immunogen at three well characterized neutralizing epitopes.

A totally different pattern of reactivity was observed with the human hybridoma, MAb 15e, prepared from an HIV-1 infected individual as described in Ho et al.; *J. Virol.* 65:489-93 (1991). With this MAb, the greatest binding was achieved with MN-rgp120 and rgp120 from C7, and the poorest reactivity was seen with the two clones of rgp120 from the C11. Moderate, but comparable reactivity was seen with rgp120s from the C10 and C17.

These results demonstrate that the 15e epitope is polymorphic, and that the epitope is conserved on MN-rgp120 and rgp120 from C7, but has been lost on rgp120s from C11. Interestingly, the two different clones of gp120 derived from C6 gave strikingly different patterns of antibody binding. Thus, rgp120 from clone C6.5 exhibited strong reactivity with this antibody, whereas rgp120 from clones C6.1 exhibited significantly weaker activity with this MAb. Comparison of sequence data (Figure 3) showed that the two C6 clones differed at 6 amino acid positions. Based on comparative binding to the other viral proteins of known sequence, it appeared that the substitution of K for I at position 351 in the C3 domain of gp120 could account for the difference in binding activity. This result is also consistent with both clones of C11 similarly containing a positively-

charged K at this position, and also being poorly reactive with this MAb. Alternatively, a T for I substitution at position 439 in the C4 domain could account for the difference in 15e binding between C6.1 and C6.5. Although the inability of the two C11 clones to bind 15e cannot be explained by polymorphism at this position in the C4 domain, they could be affected by the adjacent T for M substitution at position 434.

10 Discussion

In these studies, the viruses and immune responses in seven of nine vaccinees who became infected with HIV-1 through high risk activity while participating in Phase I or Phase 2 trials of MN-rgp120, a candidate HIV-1 vaccine were analyzed. Such infections would be expected to occur for one of two reasons: 1) lack of sufficient immune response at the time of infection; or 2) infection with viruses that fall outside of the antigenic spectrum expected to be covered by the vaccine immunogen. The data indicate that both explanations may be involved with the infections observed (Table 8).

TABLE 8
Summary of Breakthrough Infections

| 5 | <u>Case No.</u> | <u>Adequate</u> <u>Immunization</u> | MN-rgp120 <u>Homology</u> <u>(%)</u> | <u>Homologous to MN-rgp120</u> | | |
|----|-----------------|--|--|--------------------------------|-----------------------------|-----------------------------|
| | | | | <u>V3</u> <u>PND</u> | <u>C4</u> <u>Epitope</u> | <u>V2</u> <u>Epitope</u> |
| | C6 | - | 79 | + | - | - |
| | C8 | - | 78 | + | - | - |
| | C15 | - | 72 | - | - | - |
| | C7 | - | 70 | - | - | - |
| 10 | C11 | + | 75 | - | - | - |
| | C10 | + | 69 | - | - | - |
| | C17 | + | 80 | + | + | - |

15 Two of the infections occurred in individuals who failed to receive the minimum three doses of vaccine typically required for the induction of protective immunity with protein subunit vaccines (e.g. hepatitis B virus formulated in alum adjuvant as described in Francis et al.; *Ann. Int. Med.* 97:362-6 (1982)).

20 Two additional breakthrough infections occurred in vaccinees who had weak or undetectable primary (C7) and booster (C15) responses. Of the three individuals who became infected with HIV-1 after receiving three or

25 more productive immunizations (C10, C11, and C17), at least two, and possibly all three, appear to have become infected more than six months after receiving their last immunization. Because antibody titers to MN-rgp120 typically decay with a half-time of 2 to 2.5

30 months [Belshe et al.; *JAMA* 272(6):475-80 (1994); Berman et al.; *AIDS* 8:591-601 (1994)], antibody titers would be expected to have decayed at least eight-fold and possibly as much as sixty four-fold at the time of infection. Thus, the lack of a sufficient immune

response at the time of infection represents a potential explanation for at least six of the seven breakthrough infections.

5 Data from vaccine efficacy studies in gp160 immunized chimpanzees [McElrath et al.; Longitudinal Vaccine-Induced Immunity and Risk Behavior of Study Participants in AVEG Phase II Protocol 201. In: Abstracts from Eighth Annual Meeting of the National Cooperative Vaccine Development Groups for AIDS. 10 Bethesda, MD 1996:216] challenged with HIV-1, and gp120-immunized rhesus macaques challenged with a chimeric SIV/HIV-1 virus (SHIV) suggest that the magnitude of the neutralizing antibody response at the time of infection is a critical correlate of protective 15 immunity. If maintaining neutralizing antibody titers proves to be a valid correlate of protective immunity in humans, then formulations (e.g. novel adjuvants) or immunization regimes (frequent boosting) designed to maximize the antibody responses may be required to 20 achieve long lasting protection. Use of a booster every six months may be advantageous.

The other likely explanation for the late infections is the antigenic difference between the vaccine and the breakthrough virus envelope 25 glycoproteins. This explanation is supported by the observation that four of the seven breakthrough viruses possessed envelope glycoproteins that differed from the MN-rgp120 by 25-30% at the amino acid level. Differences of this magnitude have historically 30 [Myers et al.; *Retroviruses and AIDS Database*, Los Alamos National Laboratory (1992 and 1995)] been associated with inter-subtype variation and far exceeds the average 10-20% variation expected for viruses within the same subtype.

35 Although the biologic significance of sequence variation in many regions of the envelope glycoprotein

is unclear, polymorphism at neutralizing epitopes is an important factor that affects vaccine efficacy. Previous studies [Salmon-Ceron et al.; *AIDS Res. and Human Retroviruses* 11:1479-86 (1995); Javaherian et al.; *Science* 250:1590-3 (1990)] have demonstrated that the breadth of neutralizing activity that could be elicited by HIV-1 envelope derived vaccines was critically dependent on the sequence of epitopes in the V3 domain (e.g.; the PND). Thus, candidate vaccines based on the LAI strain of HIV-1 (the prototypic "non-MN-like" subtype B virus), exhibited little or no cross neutralizing activity with subtype B viruses, whereas vaccines that contained the "MN-like-" PND sequence (IGPGRAF) (SEQ. ID. NO. 44) exhibited broad cross neutralizing activity. That four of the seven breakthrough viruses possessed envelope glycoproteins with radical amino acid substitutions in the PND is consistent with the explanation that differences in antigenic structure explain some of these infections.

Over the last few years, it has become clear that polymorphism among "MN-like" viruses occurs at neutralizing epitopes outside of the PND. The best example occurs in the C4 domain where two antigenically distinct variants are distinguished by the presence of either K or E at position 429 [Moore et al.; *AIDS* 3:155-63 (1989)]. Because six of the seven breakthrough viruses differed from the vaccine strain in that they contained E rather than K at position 429, antibodies raised to the C4 domain of MN-rgp120 were unlikely to neutralize the viruses infecting in six of the seven vaccinees.

Other neutralizing epitopes are known to be present in the V1 and V2 domains of gp120. Although these regions are highly variable, due to insertions and deletions, neutralizing epitopes have been described by McKeating et al.; *J. Virol.* 67:4932-44

(1993); Moore et al.; *J. Virol.* 67:6136-6151 (1993);
and Davis et al.; *J. Gen. Virol.* 74:2609-17 (1993).
Several of these epitopes overlap an amino terminal
sequence of the V2 domain containing the tri-peptide
5 sequence RDK at positions corresponding to 142 to 144
of MN-rgp120 [McKeating et al.; *J. Virol.* 67:4932-44
(1993); Moore et al.; *J. Virol.* 67:6136-6151 (1993)].
Like the C4 epitope, variation in this sequence is
known to occur between different substrains derived
10 from the same parental isolate. Since all seven
breakthrough viruses differed from MN-rgp120 in that
they possessed the RDK sequence, rather than the GDK
sequence present in the vaccine antigen, neutralizing
antibodies to the V2 domain of MN-rgp120 would not have
15 been expected neutralize any of the viruses recovered
from the vaccinees immunized with MN-rgp120.

Although polymorphisms at neutralizing epitopes
might account for the lack of protection in most of the
infections, this does not appear to explain the
20 infection of vaccinee C17, who was infected by a virus
that matched MN-rgp120 in the V3 and C4 domains. If a
difference in sequence was responsible for the lack of
protection in this case, the critical difference might
relate to the unusual sequence in the V1 domain of
25 gp120 from this breakthrough virus. Several studies
have shown that the V1 domain possesses epitopes
recognized by virus neutralizing monoclonal antibodies
[McKeating et al.; *J. Virol.* 67:4932-44 (1993);
Davis et al.; *J. Gen. Virol.* 74:2609-17 (1993);
30 Kayman et al.; *J. Virol.* 68:400-410 (1994)].

Although far less is known about the V1 epitopes
relative to other neutralizing sites, the V1 epitopes
appear to be conformation-dependent, and antisera from
HIV-1 infected individuals recognize epitopes in the V1
35 and V2 domains [McKeating et al.; *J. Virol.* 67:4932-44
(1993); Kayman et al.; *J. Virol.* 68:400-410 (1994)].

The V1 sequence of the virus from C17 is noteworthy because it is smaller and contains fewer N-linked glycosylation sites than that of MN-rgp120 or any of the other breakthrough viruses. By the same token, the envelope glycoproteins from C11 and C6 are noteworthy because they are significantly larger and contain more glycosylation sites than MN-rgp120 or the other breakthrough viruses.

While differences in amino acid sequence can provide clues to differences in antigenic structure, the consequences of such polymorphism can only be proven through antibody binding studies. To correlate differences in sequence with differences in antigenic structure, gp120 from two clones each of all seven breakthrough viruses was expressed and the antigenicity of the clones with a panel of monoclonal antibodies was examined. As predicted from the sequence data, none of the breakthrough virus envelope glycoproteins reacted with neutralizing MAbs to the V2 domain of MN-rgp120. When MAbs to the C4 domain were examined, only the C17 envelope glycoprotein (that matched MN-rgp120 with respect to K429) showed significant, albeit lower, binding. Surprisingly, the three breakthrough envelope glycoproteins that contained the subtype B PND consensus sequence, IGPGRF (SEQ. ID. NO. !!), gave poor reactivity with all three PND directed MAbs, even though they possessed PND sequences closely related to the vaccine immunogen. Thus, all three of the vaccine isolates appeared to possess changes outside of the recognition site that interfered with MAb binding.

It has been known for many years that resistance to neutralization in vitro can sometimes be attributed to mutations in remote sequences that alter the conformation of neutralizing epitopes and interfere with recognition by virus neutralizing antibodies [Nara et al.; *J. Virol.* 64:3779-91 (1990); Cordonnier

et al.; *Nature* 340:571-4 (1989)]. Together, these results indicate that the antigenic structure of the envelope glycoproteins recovered from the breakthrough viruses differed significantly from that of the vaccine antigen.

A novel result was the localization of residues in the C3 domain that appeared to affect the binding of the virus neutralizing human MAb, 15e. This MAb is known to recognize a discontinuous epitope, block CD4 binding, and neutralize a variety of laboratory and primary isolates of HIV-1 [Ho et al.; *J. Virol.* 65:489-93 (1991); Thali et al.; *J. Virol.* 66:5635-5641 (1992); Moore et al.; *AIDS Res. Hum. Retroviruses* 9:1179-1187 (1993)].

Comparative binding to envelope glycoproteins from the breakthrough viruses indicated that recognition by this antibody is critically dependent on residues in the C3 or C4 domains of gp120. The unique occurrence of a positively charged K at position 351 in the C3 domain provides a common explanation for the inability of the C11.5, C11.7 and C6.1 strains of HIV-1 to bind to 15e. Alternatively, it is possible that different amino acid substitutions in different locations account for the failure of 15e to bind to rgp120s from the C6 and C11 clones. The only obvious positions where substitutions of this type occur are in the C4 domain where T replaces M at 434 (C11) and T replaces I at 439.

The present studies demonstrate that the current formulation of MN-rgp120 is less than 100% effective against HIV-1 infection. Based on previous in vitro and in vivo studies with MN-rgp120, protection from natural HIV-1 infection in humans is expected to depend on a threshold concentration of virus-neutralizing antibodies, and antigenic similarity between the vaccine immunogen and the challenge virus.

In this regard, only one of the seven breakthrough infections (C17) was unexpected. This individual received a full course of immunizations yet became infected with a virus similar to MN-rgp120 at at least two important neutralizing epitopes (V3 and C4 domains). This infection might be related to the magnitude of the antibody response at the time of infection, or antigenic differences between the breakthrough virus and the vaccine strain, or circumstances of infection (e.g., ulcerative lesions, infection by donor with acute infection or high viremia), not monitored in this protocol. Alternatively this individual may represent a true vaccine failure, without clear explanation.

On balance, the analysis of breakthrough infections described herein did not uncover any data that would discourage the continued development of MN-rgp120 as a vaccine to prevent HIV-1 infection. The results support speculation that enhancing vaccine immunogenicity (as by additional booster immunizations) may be required to maintain long term protective immunity, and that the addition of rgp120 from other antigenically different strains of virus in addition to MN-rgp120 are useful to expand the breadth of protection.

The availability of viruses and viral glycoproteins derived from breakthrough infections may provide an important means to streamline the process of identifying new antigens for inclusion into a multivalent vaccine. Recombinant viral glycoproteins prepared from breakthrough viruses, by definition, possess antigenic structures that are significantly different from MN-rgp120, and are be representative of viruses currently being transmitted. Thus, combining rgp120 from breakthrough viruses with MN-rgp120 is an effective way complement and significantly expand

antigenic complexity and increase breadth of cross neutralizing activity.

SEQUENCE LISTING

(1) GENERAL INFORMATION:

- (i) APPLICANT: Berman, Phillip W.
- 5 (ii) TITLE OF INVENTION: HIV ENVELOPE POLYPEPTIDES AND VACCINE
- (iii) NUMBER OF SEQUENCES: 44
- (iv) CORRESPONDENCE ADDRESS:
- 10 (A) ADDRESSEE: SKJERVEN, MORRILL, MACPHERSON, ET AL.
- (B) STREET: 25 Metro Drive, Suite 700
- (C) CITY: San Jose
- (D) STATE: California
- (E) COUNTRY: USA
- (F) ZIP: 95110
- 15 (v) COMPUTER READABLE FORM:
- (A) MEDIUM TYPE: 3.5 inch, 1.44 Mb floppy disk
- (B) COMPUTER: IBM PC compatible
- (C) OPERATING SYSTEM: PC-DOS/MS-DOS
- (D) SOFTWARE: WinPatIn (Genentech)
- 20 (vi) CURRENT APPLICATION DATA:
- (A) APPLICATION NUMBER:
- (B) FILING DATE:
- (C) CLASSIFICATION:
- 25 (viii) ATTORNEY/AGENT INFORMATION:
- (A) NAME: Terlizzi, Laura
- (B) REGISTRATION NUMBER: 31,307
- (C) REFERENCE/DOCKET NUMBER: M-3897 US
- (ix) TELECOMMUNICATION INFORMATION:
- 30 (A) TELEPHONE: (408) 453-9200
- (B) TELEFAX: (408) 453-7979

(2) INFORMATION FOR SEQ ID NO:1:

- (i) SEQUENCE CHARACTERISTICS:
- 35 (A) LENGTH: 1503 base pairs
- (B) TYPE: Nucleic Acid
- (C) STRANDEDNESS: Single
- (D) TOPOLOGY: Linear
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO:1:
- 40 GGG GTA CCT GTG TGG AAG GAA GCA ACC ACC ACT CTA 36
Gly Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu
1 5 10
- 45 TTT TCT GCA TCA GAT GCT AAA GCA TAT GAC ACA GAG GTG 75
Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val
15 20 25
- 50 CAT AAT GTT TGG GCC ACA CAT GCT TGT GTA CCC ACA GAC 114
His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp
30 35
- 55 CCA AAC CCA CAA GAA ATG GTA TTG GAA AAT GTG ACA GAA 153
Pro Asn Pro Gln Glu Met Val Leu Glu Asn Val Thr Glu
40 45 50
- GAT TTT AAC ATG TGG AAA AAT GAC ATG GTA GAA CAG ATG 192
Asp Phe Asn Met Trp Lys Asn Asp Met Val Glu Gln Met
55 60
- 60 CAT GAG GAT ATA ATC AGT TTA TGG GAT CAA AGC CTA AAA 231
His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys
65 70 75

| | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| | CCA | TGT | GTA | AAA | TTA | ACC | CCA | CTC | TGT | ATT | ACT | TTA | AAT | 270 |
| | Pro | Cys | Val | Lys | L u | Thr | Pro | Leu | Cys | Ile | Thr | Leu | Asn | |
| | | | 80 | | | | | 85 | | | | | 90 | |
| 5 | TGC | ACC | AAT | TGG | AAG | AAG | AAT | GAT | ACT | AAA | ACT | AAT | AGT | 309 |
| | Cys | Thr | Asn | Trp | Lys | Lys | Asn | Asp | Thr | Lys | Thr | Asn | Ser | |
| | | | | 95 | | | | 100 | | | | | | |
| 10 | AGT | AGT | ACT | ACA | ACT | AAT | AAT | AGT | AGT | GCT | ACA | GCT | AAT | 348 |
| | Ser | Ser | Thr | Thr | Thr | Asn | Asn | Ser | Ser | Ala | Thr | Ala | Asn | |
| | | | 105 | | | | 110 | | | | | 115 | | |
| 15 | AGT | AGT | AGT | ACT | ACA | ACT | AAT | AGT | AGT | TGG | GGA | GAG | ATA | 387 |
| | Ser | Ser | Ser | Thr | Thr | Thr | Asn | Ser | Ser | Trp | Gly | Glu | Ile | |
| | | | | 120 | | | | 125 | | | | | | |
| 20 | AAG | GAG | GGA | GAA | ATA | AAG | AAC | TGC | TCT | TTC | AAT | ATC | ACC | 426 |
| | Lys | Glu | Gly | Glu | Ile | Lys | Asn | Cys | Ser | Phe | Asn | Ile | Thr | |
| | | 130 | | | | 135 | | | | | 140 | | | |
| | ACA | AGC | ATA | AGA | GAC | AAG | GTG | AAG | AAA | GAA | TAT | GCA | CTT | 465 |
| | Thr | Ser | Ile | Arg | Asp | Lys | Val | Lys | Lys | Glu | Tyr | Ala | Leu | |
| | | | 145 | | | | 150 | | | | | | 155 | |
| 25 | TTT | TAT | AGC | CTT | GAT | GTA | GTA | CCA | ATA | GAA | AAT | GAT | AAT | 504 |
| | Phe | Tyr | Ser | Leu | Asp | Val | Val | Pro | Ile | Glu | Asn | Asp | Asn | |
| | | | | | 160 | | | | | 165 | | | | |
| 30 | ACT | AGC | TAT | AGG | TTG | AGA | AGT | TGT | AAC | ACC | TCA | GTC | ATT | 543 |
| | Thr | Ser | Tyr | Arg | Leu | Arg | Ser | Cys | Asn | Thr | Ser | Val | Ile | |
| | | | 170 | | | | 175 | | | | | 180 | | |
| 35 | ACA | CAA | GCC | TGT | CCA | AAG | GTA | ACT | TTT | GAG | CCA | ATT | CCC | 582 |
| | Thr | Gln | Ala | Cys | Pro | Lys | Val | Thr | Phe | Glu | Pro | Ile | Pro | |
| | | | | 185 | | | | | 190 | | | | | |
| 40 | ATA | CAT | TAT | TGT | ACC | CCG | GCT | GGT | TTT | GCG | ATT | CTG | AAG | 621 |
| | Ile | His | Tyr | Cys | Thr | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | |
| | | 195 | | | | 200 | | | | | 205 | | | |
| | TGT | AGA | GAT | AAA | AAG | TTC | AAT | GGA | ACA | GGA | CCA | TGC | AAA | 660 |
| | Cys | Arg | Asp | Lys | Lys | Phe | Asn | Gly | Thr | Gly | Pro | Cys | Lys | |
| | | | 210 | | | | 215 | | | | | 220 | | |
| 45 | AAT | GTT | AGC | ACA | GTA | CAA | TGT | GCA | CAT | GGA | ATT | AAG | CCA | 699 |
| | Asn | Val | Ser | Thr | Val | Gln | Cys | Ala | His | Gly | Ile | Lys | Pro | |
| | | | | | 225 | | | | 230 | | | | | |
| 50 | GTA | GTG | TCA | ACT | CAA | CTG | CTG | TTA | AAT | GGC | AGC | CTA | GCA | 738 |
| | Val | Val | Ser | Thr | Gln | Leu | Leu | Leu | Asn | Gly | Ser | Leu | Ala | |
| | | 235 | | | | 240 | | | | | | 245 | | |
| 55 | GAA | GAA | GAG | GTA | ATA | ATT | AGA | TCT | GCC | AAT | TTC | TCA | AAC | 777 |
| | Glu | Glu | Glu | Val | Ile | Ile | Arg | Ser | Ala | Asn | Phe | Ser | Asn | |
| | | | | 250 | | | | 255 | | | | | | |
| 60 | AAT | GCT | AAA | ATC | ATA | ATA | GTA | CAG | TTG | AGG | GAA | CCT | GTA | 816 |
| | Asn | Ala | Lys | Ile | Ile | Ile | Val | Gln | Leu | Arg | Glu | Pro | Val | |
| | | 260 | | | | 265 | | | | | 270 | | | |
| | GAA | ATT | AAT | TGT | ACA | AGA | CCC | AGC | AAC | AAT | ACA | ATA | AAA | 855 |
| | Glu | Ile | Asn | Cys | Thr | Arg | Pro | Ser | Asn | Asn | Thr | Ile | Lys | |
| | | | 275 | | | | | 280 | | | | | 285 | |

GGT ATA CAC ATA GGA CCA GGG AGA GCA TTT TAT GCA ACA 894
 Gly Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala Thr
 290 295

5 GGA GAC ATA CGA GGA GAT ATA AGA CAA GCA CAT TGT AAC 933
 Gly Asp Ile Arg Gly Asp Ile Arg Gln Ala His Cys Asn
 300 305 310

10 ATT AGT GGA GCA AAA TGG AAT AAC ACT TTA AAG AAG GTA 972
 Ile Ser Gly Ala Lys Trp Asn Asn Thr Leu Lys Lys Val
 315 320

15 GTT AAA AAA TTA AAA GAA CAA TTT CCA AAT AAA ACA ATA 1011
 Val Lys Lys Leu Lys Glu Gln Phe Pro Asn Lys Thr Ile
 325 330 335

GTC TTT AAC CAT TCC TCA GGA GGG GAC CCA GAA ATT GTA 1050
 Val Phe Asn His Ser Ser Gly Gly Asp Pro Glu Ile Val
 340 345 350

20 ATG CAC AGT TTT AAT TGT CAA GGG GAA TTT TTC TAC TGT 1089
 Met His Ser Phe Asn Cys Gln Gly Glu Phe Phe Tyr Cys
 355 360

25 AAT ACA ACA AAG CTG TTT AAT AGT ACT TGG AAT GAT ACT 1128
 Asn Thr Thr Lys Leu Phe Asn Ser Thr Trp Asn Asp Thr
 365 370 375

30 ACA GAG TCA AAT AAC AAT GAT AGT ACT ATT ACA CTC CCA 1167
 Thr Glu Ser Asn Asn Asn Asp Ser Thr Ile Thr Leu Pro
 380 385

35 TGC AGA ATA AAA CAA ATT ATA AAC ATG TGG CAG GAA ATA 1206
 Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Ile
 390 395 400

GGA AAA GCA ATG TAT GCC CCT CCC ACC AGA GGA GAA ATT 1245
 Gly Lys Ala Met Tyr Ala Pro Pro Thr Arg Gly Glu Ile
 405 410 415

40 AAA TGT TCA TCA AAT ATT ACA GGA CTA CTG TTA ATA AGA 1284
 Lys Cys Ser Ser Asn Ile Thr Gly Leu Leu Ile Arg
 420 425

45 GAT GGT GGT ATT AAC ACT AGC GAT GCC ACC GAG ACC TTC 1323
 Asp Gly Gly Ile Asn Thr Ser Asp Ala Thr Glu Thr Phe
 430 435 440

50 AGA CCG GGA GGA GGA GAT ATG AGG GAC AAT TGG AGA AGT 1362
 Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser
 445 450

55 GAA TTA TAT AAA TAT AAA GTA GTG AAA ATT GAG CCA TTA 1401
 Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu
 455 460 465

GGA GTA GCA CCC ACC AAG GCA AAG AGA AGA GTG GTG CAG 1440
 Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 470 475 480

60 AGA GAA AAA AGA GCA GTA ACA CTA GGA GCT ATG TTC CTT 1479
 Arg Glu Lys Arg Ala Val Thr Leu Gly Ala Met Phe Leu
 485 490

GGG TTC TTA GGA GCA TAA AGC TTC 1503
 Gly Phe Leu Gly Ala Xaa Ser Phe
 495 500 501

5 (2) INFORMATION FOR SEQ ID NO:2:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 501 amino acids

(B) TYPE: Amino Acid

(D) TOPOLOGY: Linear

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:2:

| | | | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|--|
| | Gly | Val | Pro | Val | Trp | Lys | Glu | Ala | Thr | Thr | Thr | Leu | Phe | Cys | Ala | |
| | 1 | | | | 5 | | | | | 10 | | | | | 15 | |
| 15 | Ser | Asp | Ala | Lys | Ala | Tyr | Asp | Thr | Glu | Val | His | Asn | Val | Trp | Ala | |
| | | | | 20 | | | | | | 25 | | | | | 30 | |
| | Thr | His | Ala | Cys | Val | Pro | Thr | Asp | Pro | Asn | Pro | Gln | Glu | Met | Val | |
| | | | | 35 | | | | | | 40 | | | | | 45 | |
| 20 | Leu | Glu | Asn | Val | Thr | Glu | Asp | Phe | Asn | Met | Trp | Lys | Asn | Asp | Met | |
| | | | | 50 | | | | | | 55 | | | | | 60 | |
| | Val | Glu | Gln | Met | His | Glu | Asp | Ile | Ile | Ser | Leu | Trp | Asp | Gln | Ser | |
| 25 | | | | 65 | | | | | | 70 | | | | | 75 | |
| | Leu | Lys | Pro | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Ile | Thr | Leu | Asn | |
| | | | | 80 | | | | | | 85 | | | | | 90 | |
| 30 | Cys | Thr | Asn | Trp | Lys | Lys | Asn | Asp | Thr | Lys | Thr | Asn | Ser | Ser | Ser | |
| | | | | 95 | | | | | | 100 | | | | | 105 | |
| | Thr | Thr | Thr | Asn | Asn | Ser | Ser | Ala | Thr | Ala | Asn | Ser | Ser | Ser | Thr | |
| | | | | 110 | | | | | | 115 | | | | | 120 | |
| 35 | Thr | Thr | Asn | Ser | Ser | Trp | Gly | Glu | Ile | Lys | Glu | Gly | Glu | Ile | Lys | |
| | | | | 125 | | | | | | 130 | | | | | 135 | |
| | Asn | Cys | Ser | Phe | Asn | Ile | Thr | Thr | Ser | Ile | Arg | Asp | Lys | Val | Lys | |
| 40 | | | | 140 | | | | | | 145 | | | | | 150 | |
| | Lys | Glu | Tyr | Ala | Leu | Phe | Tyr | Ser | Leu | Asp | Val | Val | Pro | Ile | Glu | |
| | | | | 155 | | | | | | 160 | | | | | 165 | |
| 45 | Asn | Asp | Asn | Thr | Ser | Tyr | Arg | Leu | Arg | Ser | Cys | Asn | Thr | Ser | Val | |
| | | | | 170 | | | | | | 175 | | | | | 180 | |
| | Ile | Thr | Gln | Ala | Cys | Pro | Lys | Val | Thr | Phe | Glu | Pro | Ile | Pro | Ile | |
| | | | | 185 | | | | | | 190 | | | | | 195 | |
| 50 | His | Tyr | Cys | Thr | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Arg | Asp | |
| | | | | 200 | | | | | | 205 | | | | | 210 | |
| | Lys | Lys | Phe | Asn | Gly | Thr | Gly | Pro | Cys | Lys | Asn | Val | Ser | Thr | Val | |
| 55 | | | | 215 | | | | | | 220 | | | | | 225 | |
| | Gln | Cys | Ala | His | Gly | Ile | Lys | Pro | Val | Val | Ser | Thr | Gln | Leu | Leu | |
| | | | | 230 | | | | | | 235 | | | | | 240 | |
| 60 | Leu | Asn | Gly | Ser | Leu | Ala | Glu | Glu | Glu | Val | Ile | Ile | Arg | Ser | Ala | |
| | | | | 245 | | | | | | 250 | | | | | 255 | |
| | Asn | Phe | Ser | Asn | Asn | Ala | Lys | Ile | Ile | Ile | Val | Gln | Leu | Arg | Glu | |
| 65 | | | | 260 | | | | | | 265 | | | | | 270 | |

Pro Val Glu Ile Asn Cys Thr Arg Pro Ser Asn Asn Thr Ile Lys
 275 280 285
 5 Gly Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala Thr Gly Asp
 290 295 300
 Ile Arg Gly Asp Ile Arg Gln Ala His Cys Asn Ile Ser Gly Ala
 305 310 315
 10 Lys Trp Asn Asn Thr Leu Lys Lys Val Val Lys Lys Leu Lys Glu
 320 325 330
 Gln Phe Pro Asn Lys Thr Ile Val Phe Asn His Ser Ser Gly Gly
 335 340 345
 15 Asp Pro Glu Ile Val Met His Ser Phe Asn Cys Gln Gly Glu Phe
 350 355 360
 Phe Tyr Cys Asn Thr Thr Lys Leu Phe Asn Ser Thr Trp Asn Asp
 365 370 375
 20 Thr Thr Glu Ser Asn Asn Asn Asp Ser Thr Ile Thr Leu Pro Cys
 380 385 390
 25 Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Ile Gly Lys Ala
 395 400 405
 Met Tyr Ala Pro Pro Thr Arg Gly Glu Ile Lys Cys Ser Ser Asn
 410 415 420
 30 Ile Thr Gly Leu Leu Leu Ile Arg Asp Gly Gly Ile Asn Thr Ser
 425 430 435
 Asp Ala Thr Glu Thr Phe Arg Pro Gly Gly Gly Asp Met Arg Asp
 440 445 450
 35 Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu
 455 460 465
 40 Pro Leu Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 470 475 480
 Arg Glu Lys Arg Ala Val Thr Leu Gly Ala Met Phe Leu Gly Phe
 485 490 495
 45 Leu Gly Ala Xaa Ser Phe
 500 501

(2) INFORMATION FOR SEQ ID NO:3:

- 50 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1503 base pairs
 (B) TYPE: Nucleic Acid
 (C) STRANDEDNESS: Single
 (D) TOPOLOGY: Linear
 55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:3:

GGG GTA CCT GTA TGG AAA GAA GCA ACC ACC ACT CTA 36
 Gly Val Pro Val Trp Lys Glu Ala Thr Thr Leu
 1 5 10
 60 TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GAG GTG 75
 Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val
 15 20 25

| | | |
|----|---|-----|
| | CAT AAT GTT TGG GCC ACA CAT GCT TGT GTA CCC ACA GAC | 114 |
| | His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp | |
| | 30 35 | |
| 5 | CCA AAC CCA CAA GAA ATG GTA TTG GAA AAT GTG ACA GAA | 153 |
| | Pro Asn Pro Gln Glu Met Val Leu Glu Asn Val Thr Glu | |
| | 40 45 50 | |
| 10 | GAT TTT AAC ATG TGG AAA AAT GAC ATG GTA GAA CAG ATG | 192 |
| | Asp Phe Asn Met Trp Lys Asn Asp Met Val Glu Gln Met | |
| | 55 60 | |
| 15 | CAT GAG ANT ATA ATC AGT TTA TGG GAT CAA AGC CTA AAA | 231 |
| | His Glu Xaa Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys | |
| | 65 70 75 | |
| 20 | CCA TGT GTA AAA TTA ACC CCA CTC TGT ATT ACT TTA AAT | 270 |
| | Pro Cys Val Lys Leu Thr Pro Leu Cys Ile Thr Leu Asn | |
| | 80 85 90 | |
| | TGC ACC AAT TGG AAG GAG AAT GAT ACT AAA ACT AAT AGT | 309 |
| | Cys Thr Asn Trp Lys Glu Asn Asp Thr Lys Thr Asn Ser | |
| | 95 100 | |
| 25 | AGT AGT ACT ACA ACT AAT AAT AGT AGT GCT ACA GCT AAT | 348 |
| | Ser Ser Thr Thr Thr Asn Asn Ser Ser Ala Thr Ala Asn | |
| | 105 110 115 | |
| 30 | AGT AGT ACT ACT ACA ACT AAT AGT AGT TGG GGA GAC ATA | 387 |
| | Ser Ser Ser Thr Thr Thr Asn Ser Ser Trp Gly Glu Ile | |
| | 120 125 | |
| 35 | AAG GAG GGA GAA ATA AAG AAC TGC TCT TTC AAT ATC ACC | 426 |
| | Lys Glu Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile Thr | |
| | 130 135 140 | |
| 40 | ACA GGC ATA AGA GAC AAG GTG AAG AAA GAA TAT GCA CTT | 465 |
| | Thr Gly Ile Arg Asp Lys Val Lys Lys Glu Tyr Ala Leu | |
| | 145 150 155 | |
| | TTT TAT AGC CTT GAT GTA GTA CCA ATA GAA AAT GAT AAT | 504 |
| | Phe Tyr Ser Leu Asp Val Val Pro Ile Glu Asn Asp Asn | |
| | 160 165 | |
| 45 | ACT AGC TAT AGG TTG AGA AGT TGT AAC ACC TCA GTC ATT | 543 |
| | Thr Ser Tyr Arg Leu Arg Ser Cys Asn Thr Ser Val Ile | |
| | 170 175 180 | |
| 50 | ACA CAA GCC TGT CCA AAG GTA ACT TTT GAG CCA ATT CCC | 582 |
| | Thr Gln Ala Cys Pro Lys Val Thr Phe Glu Pro Ile Pro | |
| | 185 190 | |
| 55 | ATA CAT TAT TGT ACC CCG GCT GGT TTT GCG ATT CTG AAG | 621 |
| | Ile His Tyr Cys Thr Pro Ala Gly Phe Ala Ile Leu Lys | |
| | 195 200 205 | |
| 60 | TGT AAA GAT AAA AAG TTC AAT GGA ACA GGA CCA TGC AAA | 660 |
| | Cys Lys Asp Lys Lys Phe Asn Gly Thr Gly Pro Cys Lys | |
| | 210 215 220 | |
| | AAT GTT AGC ACA GTA CAA TGT ACA CAT GGA ATT AAG CCA | 699 |
| | Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Lys Pro | |
| | 225 230 | |

-97-

AGA CCG GGA GGA GGA GAT ATG AGG GAC AAT TGG AGA AGT 1362
 Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser
 445 450

5 GAA TTA TAT AAA TAT AAA GTA GTG AAA ATT GAG CCA TTA 1401
 Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu
 455 460 465

10 GGA GTA GCA CCC ACC AAG GCA AAG AGA AGA GTG GTG CAG 1440
 Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln
 470 475 480

15 AGA GAA AAA AGA GCA GTA ACA CTA GGA GCT ATG TTC CTT 1479
 Arg Glu Lys Arg Ala Val Thr Leu Gly Ala Met Phe Leu
 485 490

GGG TTC TTG GGA GCA TAA AGC TTC 1503
 Gly Phe Leu Gly Ala Xaa Ser Phe
 495 500 501

20 (2) INFORMATION FOR SEQ ID NO:4:
 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 501 amino acids
 (B) TYPE: Amino Acid
 (D) TOPOLOGY: Linear

25 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:4:

Gly Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe Cys Ala
 1 5 10 15

30 Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val His Asn Val Trp Ala
 20 25 30

35 Thr His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Met Val
 35 40 45

Leu Glu Asn Val Thr Glu Asp Phe Asn Met Trp Lys Asn Asp Met
 50 55 60

40 Val Glu Gln Met His Glu Xaa Ile Ile Ser Leu Trp Asp Gln Ser
 65 70 75

Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Ile Thr Leu Asn
 80 85 90

45 Cys Thr Asn Trp Lys Glu Asn Asp Thr Lys Thr Asn Ser Ser Ser
 95 100 105

50 Thr Thr Thr Asn Asn Ser Ser Ala Thr Ala Asn Ser Ser Ser Thr
 110 115 120

Thr Thr Asn Ser Ser Trp Gly Glu Ile Lys Glu Gly Glu Ile Lys
 125 130 135

55 Asn Cys Ser Phe Asn Ile Thr Thr Gly Ile Arg Asp Lys Val Lys
 140 145 150

Lys Glu Tyr Ala Leu Phe Tyr Ser Leu Asp Val Val Pro Ile Glu
 155 160 165

60 Asn Asp Asn Thr Ser Tyr Arg Leu Arg Ser Cys Asn Thr Ser Val
 170 175 180

65 Ile Thr Gln Ala Cys Pro Lys Val Thr Phe Glu Pro Ile Pro Ile
 185 190 195

| | | | | |
|----|-----------------|---------------------|---------------------|-----|
| | His Tyr Cys Thr | Pro Ala Gly Phe Ala | Ile Leu Lys Cys Lys | Asp |
| | | 200 | 205 | 210 |
| 5 | Lys Lys Phe Asn | Gly Thr Gly Pro Cys | Lys Asn Val Ser Thr | Val |
| | | 215 | 220 | 225 |
| | Gln Cys Thr His | Gly Ile Lys Pro Val | Val Ser Thr Gln Leu | Leu |
| | | 230 | 235 | 240 |
| 10 | Leu Asn Gly Ser | Leu Ala Glu Glu Glu | Val Ile Ile Arg Ser | Ala |
| | | 245 | 250 | 255 |
| | Asn Phe Ser Asn | Asn Ala Lys Ile Ile | Ile Val Gln Leu Lys | Glu |
| | | 260 | 265 | 270 |
| 15 | Pro Val Glu Ile | Asn Cys Thr Arg Pro | Ser Asn Asn Thr Ile | Lys |
| | | 275 | 280 | 285 |
| 20 | Gly Ile His Ile | Gly Pro Gly Arg Ala | Phe Tyr Ala Thr Gly | Asp |
| | | 290 | 295 | 300 |
| | Ile Arg Gly Asp | Ile Arg Gln Ala His | Cys Asn Ile Ser Gly | Ala |
| | | 305 | 310 | 315 |
| 25 | Lys Trp Asn Asn | Thr Leu Lys Lys Val | Val Ile Lys Leu Lys | Glu |
| | | 320 | 325 | 330 |
| | Gln Phe Pro Asn | Lys Thr Ile Val Phe | Asn His Ser Ser Gly | Gly |
| | | 335 | 340 | 345 |
| 30 | Asp Pro Glu Ile | Val Met His Ser Phe | Asn Cys Gln Gly Glu | Phe |
| | | 350 | 355 | 360 |
| 35 | Phe Tyr Cys Asn | Thr Thr Lys Leu Phe | Asn Ser Thr Trp Asn | Asp |
| | | 365 | 370 | 375 |
| | Thr Thr Glu Ser | Asn Asn Asn Asp Ser | Thr Ile Thr Leu Pro | Cys |
| | | 380 | 385 | 390 |
| 40 | Arg Ile Lys Gln | Ile Ile Asn Met Trp | Gln Glu Val Gly Lys | Ala |
| | | 395 | 400 | 405 |
| | Met Tyr Ala Pro | Pro Ile Arg Gly Glu | Ile Lys Cys Ser Ser | Asn |
| | | 410 | 415 | 420 |
| 45 | Ile Thr Gly Leu | Leu Leu Thr Arg Asp | Gly Gly Ile Asn Thr | Ser |
| | | 425 | 430 | 435 |
| 50 | Asp Ala Thr Glu | Thr Phe Arg Pro Gly | Gly Gly Asp Met Arg | Asp |
| | | 440 | 445 | 450 |
| | Asn Trp Arg Ser | Glu Leu Tyr Lys Tyr | Lys Val Val Lys Ile | Glu |
| | | 455 | 460 | 465 |
| 55 | Pro Leu Gly Val | Ala Pro Thr Lys Ala | Lys Arg Arg Val Val | Gln |
| | | 470 | 475 | 480 |
| | Arg Glu Lys Arg | Ala Val Thr Leu Gly | Ala Met Phe Leu Gly | Phe |
| | | 485 | 490 | 495 |
| 60 | Leu Gly Ala Xaa | Ser Phe | | |
| | | 500 501 | | |

65 (2) INFORMATION FOR SEQ ID NO:5:
(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1461 base pairs

(B) TYPE: Nucleic Acid

(C) STRANDEDNESS: Single

(D) TOPOLOGY: Linear

5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:5:

G GTA CCT GTA TGG AAA GAA GCA ACC ACC ACT CTA TTT 37
 Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe
 1 5 10

10 TGT GCA TCA GAT GCT AAA GCA TAT GAT ACA GAG GTA CAT 76
 Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val His
 15 20 25

15 AAT GTT TGG GCT ACA CAT GCC TGT GTA CCC ACA GAC CCC 115
 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro
 30 35

20 AAC CCA CAA GAA GTA GTA TTG GAA AAT GTA ACA GAA AAT 154
 Asn Pro Gln Glu Val Val Leu Glu Asn Val Thr Glu Asn
 40 45 50

25 TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAG ATG CAT 193
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His
 55 60

30 GAG GAT ATA ATC AGT TTA TGG GAT CAA AGT CTA AAG CCA 232
 Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys Pro
 65 70 75

35 TGT GTA AAA TTA ACC CCA CTC TGT GTT ACT TTA AAT TCC 271
 Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys
 80 85 90

40 ACT AAT TTG GAG AAT GCT AAT AAT ACC GAG AAT GCT AAT 310
 Thr Asn Leu Glu Asn Ala Asn Asn Thr Glu Asn Ala Asn
 95 100

45 AAT ACC AAT AAT TAT ACC TTG GGG ATG GAG AGA GGT GAA 349
 Asn Thr Asn Asn Tyr Thr Leu Gly Met Glu Arg Gly Glu
 105 110 115

50 ATA AAA AAC TGC TCT TTC AAT ATC ACC ACA AGC TTA AGA 388
 Ile Lys Asn Cys Ser Phe Asn Ile Thr Thr Ser Leu Arg
 120 125

55 GAT AAG GTG AAA AAA GAA TAT GCA TTG TTT TAT AAA CTT 427
 Asp Lys Val Lys Lys Glu Tyr Ala Leu Phe Tyr Lys Leu
 130 135 140

60 GAT GTA GTA CAA ATA GAT AAT AGT ACC AAC TAT AGG CTG 466
 Asp Val Val Gln Ile Asp Asn Ser Thr Asn Tyr Arg Leu
 145 150 155

65 ATA AGT TGT AAT ACC TCA GTC ATT ACA CAG GCC TGT CCA 505
 Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro
 160 165

70 AAG GTA TCC TTT GAG CTA ATT CCC ATA CAT TAT TGT GCC 544
 Lys Val Ser Phe Glu Leu Ile Pro Ile His Tyr Cys Ala
 170 175 180

75 CCG GCT GGT TTT GCG ATT CTA AAG TGT AAA GAT AAG AAG 583
 Pro Ala Gly Phe Ala Ile Leu Lys Cys Lys Asp Lys Lys
 185 190

TTC AAT GGA ACA GGA CCA TGT AAA AAT GTC AGC ACA GTA 622
 Phe Asn Gly Thr Gly Pro Cys Lys Asn Val Ser Thr Val
 195 200 205

5 CAA TGT ACA CAT GGA ATT AGA CCA GTA GTA TCA ACT CAA 661
 Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr Gln
 210 215 220

10 CTA CTG TTA AAT GGC AGT CTA GCA GAA GAA GAG ATA GTA 700
 Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Ile Val
 225 230

15 ATT AGA TCT GAA AAT ATC ACA GAC AAT GCT AAA ACC ATA 739
 Ile Arg Ser Glu Asn Ile Thr Asp Asn Ala Lys Thr Ile
 235 240 245

20 ATA GTG CAG CTA AAT GAA TCT ATA GTG ATT AAT TGT ACA 778
 Ile Val Gln Leu Asn Glu Ser Ile Val Ile Asn Cys Thr
 250 255

25 AGA CCC AAT AAC AAC ACA AGA AAA AGT ATA AAT ATA GGA 817
 Arg Pro Asn Asn Asn Thr Arg Lys Ser Ile Asn Ile Gly
 260 265 270

30 CCA GGG AGA GCA TTC TAT ACA ACA GGA GAC ATA ATA GGA 856
 Pro Gly Arg Ala Phe Tyr Thr Thr Gly Asp Ile Ile Gly
 275 280 285

35 GAT ATA AGA CAA GCA CAT TGT AAC CTT AGT AAA ACA CAA 895
 Asp Ile Arg Gln Ala His Cys Asn Leu Ser Lys Thr Gln
 290 295

40 TGG GAA AAA ACG TTA AGA CAG ATA GCT ATA AAA TTA GAA 934
 Trp Glu Lys Thr Leu Arg Gln Ile Ala Ile Lys Leu Glu
 300 305 310

45 GAA AAA TTT AAG AAT AAA ACA ATA GCC TTT AAT AAA TCC 973
 Glu Lys Phe Lys Asn Lys Thr Ile Ala Phe Asn Lys Ser
 315 320

50 TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC AGT TTT AAT 1012
 Ser Gly Gly Asp Pro Glu Ile Val Met His Ser Phe Asn
 325 330 335

55 TGT GGA GGG GAA TTT TTC TAC TGT AAT ACA ACA AAA CTG 1051
 Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr Lys Leu
 340 345 350

60 TTT AAT AGT ACC TGG AAT TTA ACA CAA CCG TTT AGT AAT 1090
 Phe Asn Ser Thr Trp Asn Leu Thr Gln Pro Phe Ser Asn
 355 360

ACC GGG AAT CGT ACT GAA GAG TTA AAT ATT ACA CTC CCA 1129
 Thr Gly Asn Arg Thr Glu Glu Leu Asn Ile Thr Leu Pro
 365 370 375

TGC AGA ATA AAA CAA ATC ATA AAC TTG TGG CAG GAA GTA 1168
 Cys Arg Ile Lys Gln Ile Ile Asn Leu Trp Gln Glu Val
 380 385

GGC AAA GCA ATG TAT GCC CCT CCC ATC AGA GGA CAA ATT 1207
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile
 390 395 400

AGA TGT TCA TCA AAT ATT ACA GGG CTA CTA TTA ACA AGA 1246
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg
 405 410 415

5 GAT GGT GGA AGT AAC ACC GGT GAC AAC AGG ACT GAG ACC 1285
 Asp Gly Gly Ser Asn Thr Gly Asp Asn Arg Thr Glu Thr
 420 425

10 TTT AGA CCT GGA GGA GGA GAT ATG AGG GAC AAT TGG AGA 1324
 Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 430 435 440

15 AGT GAA TTA TAT AAA TAT AAA GTA GTA AGA ATT GAA CCA 1363
 Ser Glu Leu Tyr Lys Tyr Lys Val Val Arg Ile Glu Pro
 445 450

20 TTA GGA GTA GCA CCC ACC CAG GCA AAG AGA AGA GTG GTG 1402
 Leu Gly Val Ala Pro Thr Gln Ala Lys Arg Arg Val Val
 455 460 465

CAA ACA GAA AAA AGA GCA GTG GGG ATA GGA GCT ATG TTC 1441
 Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Met Phe
 470 475 480

25 CTT GGG TTC TTG GGA GAT AA 1461
 Leu Gly Phe Leu Gly Asp
 485 486

(2) INFORMATION FOR SEQ ID NO:6:
 30 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 486 amino acids
 (B) TYPE: Amino Acid
 (D) TOPOLOGY: Linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:6:

35 Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe Cys Ala Ser
 1 5 10 15

40 Asp Ala Lys Ala Tyr Asp Thr Glu Val His Asn Val Trp Ala Thr
 20 25 30

His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Val Val Leu
 35 40 45

45 Glu Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val
 50 55 60

Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu
 65 70 75

50 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys
 80 85 90

55 Thr Asn Leu Glu Asn Ala Asn Asn Thr Glu Asn Ala Asn Asn Thr
 95 100 105

Asn Asn Tyr Thr Leu Gly Met Glu Arg Gly Glu Ile Lys Asn Cys
 110 115 120

60 Ser Phe Asn Ile Thr Thr Ser Leu Arg Asp Lys Val Lys Lys Glu
 125 130 135

Tyr Ala Leu Phe Tyr Lys Leu Asp Val Val Gln Ile Asp Asn Ser
 140 145 150

65

| | | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| | Thr | Asn | Tyr | Arg | Leu | Ile | Ser | Cys | Asn | Thr | Ser | Val | Ile | Thr | Gln |
| | | | | | 155 | | | | | 160 | | | | | 165 |
| 5 | Ala | Cys | Pro | Lys | Val | Ser | Phe | Glu | Leu | Ile | Pro | Ile | His | Tyr | Cys |
| | | | | | 170 | | | | | 175 | | | | | 180 |
| | Ala | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Lys | Asp | Lys | Lys | Phe |
| | | | | | 185 | | | | | 190 | | | | | 195 |
| 10 | Asn | Gly | Thr | Gly | Pro | Cys | Lys | Asn | Val | Ser | Thr | Val | Gln | Cys | Thr |
| | | | | | 200 | | | | | 205 | | | | | 210 |
| | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | Gln | Leu | Leu | Leu | Asn | Gly |
| | | | | | 215 | | | | | 220 | | | | | 225 |
| 15 | Ser | Leu | Ala | Glu | Glu | Glu | Ile | Val | Ile | Arg | Ser | Glu | Asn | Ile | Thr |
| | | | | | 230 | | | | | 235 | | | | | 240 |
| | Asp | Asn | Ala | Lys | Thr | Ile | Ile | Val | Gln | Leu | Asn | Glu | Ser | Ile | Val |
| | | | | | 245 | | | | | 250 | | | | | 255 |
| 20 | Ile | Asn | Cys | Thr | Arg | Pro | Asn | Asn | Asn | Thr | Arg | Lys | Ser | Ile | Asn |
| | | | | | 260 | | | | | 265 | | | | | 270 |
| | Ile | Gly | Pro | Gly | Arg | Ala | Phe | Tyr | Thr | Thr | Gly | Asp | Ile | Ile | Gly |
| | | | | | 275 | | | | | 280 | | | | | 285 |
| | Asp | Ile | Arg | Gln | Ala | His | Cys | Asn | Leu | Ser | Lys | Thr | Gln | Trp | Glu |
| | | | | | 290 | | | | | 295 | | | | | 300 |
| 30 | Lys | Thr | Leu | Arg | Gln | Ile | Ala | Ile | Lys | Leu | Glu | Glu | Lys | Phe | Lys |
| | | | | | 305 | | | | | 310 | | | | | 315 |
| | Asn | Lys | Thr | Ile | Ala | Phe | Asn | Lys | Ser | Ser | Gly | Gly | Asp | Pro | Glu |
| | | | | | 320 | | | | | 325 | | | | | 330 |
| | Ile | Val | Met | His | Ser | Phe | Asn | Cys | Gly | Gly | Glu | Phe | Phe | Tyr | Cys |
| | | | | | 335 | | | | | 340 | | | | | 345 |
| 40 | Asn | Thr | Thr | Lys | Leu | Phe | Asn | Ser | Thr | Trp | Asn | Leu | Thr | Gln | Pro |
| | | | | | 350 | | | | | 355 | | | | | 360 |
| | Phe | Ser | Asn | Thr | Gly | Asn | Arg | Thr | Glu | Glu | Leu | Asn | Ile | Thr | Leu |
| | | | | | 365 | | | | | 370 | | | | | 375 |
| 45 | Pro | Cys | Arg | Ile | Lys | Gln | Ile | Ile | Asn | Leu | Trp | Gln | Glu | Val | Gly |
| | | | | | 380 | | | | | 385 | | | | | 390 |
| | Lys | Ala | Met | Tyr | Ala | Pro | Pro | Ile | Arg | Gly | Gln | Ile | Arg | Cys | Ser |
| | | | | | 395 | | | | | 400 | | | | | 405 |
| | Ser | Asn | Ile | Thr | Gly | Leu | Leu | Leu | Thr | Arg | Asp | Gly | Gly | Ser | Asn |
| | | | | | 410 | | | | | 415 | | | | | 420 |
| 55 | Thr | Gly | Asp | Asn | Arg | Thr | Glu | Thr | Phe | Arg | Pro | Gly | Gly | Gly | Asp |
| | | | | | 425 | | | | | 430 | | | | | 435 |
| | Met | Arg | Asp | Asn | Trp | Arg | Ser | Glu | Leu | Tyr | Lys | Tyr | Lys | Val | Val |
| | | | | | 440 | | | | | 445 | | | | | 450 |
| 60 | Arg | Ile | Glu | Pro | Leu | Gly | Val | Ala | Pro | Thr | Gln | Ala | Lys | Arg | Arg |
| | | | | | 455 | | | | | 460 | | | | | 465 |
| | Val | Val | Gln | Arg | Glu | Lys | Arg | Ala | Val | Gly | Ile | Gly | Ala | Met | Phe |
| | | | | | 470 | | | | | 475 | | | | | 480 |

Leu Gly Phe Leu Gly Asp
485 486

(2) INFORMATION FOR SEQ ID NO:7:

5

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 1474 base pairs
(B) TYPE: Nucleic Acid
(C) STRANDEDNESS: Single
(D) TOPOLOGY: Linear

10

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:7:

G GTA CCT GTG TGG AAA GAA GCA ACC ACC ACT CTA TTT 37
Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe
1 5 10

TGT GCA TCA GAT GCT AAA GCA TAT GAT ACA GAG GTA CAT 76
Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val His
15 20 25

AAT GTT TGG GCT ACA CAT GCC TGT GTA CCC ACA GAC CCC 115
Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro
30 35

AAC CCA CAA GAA GTA GTA TTG GAA AAT GTA ACA GAA AAT 154
Asn Pro Gln Glu Val Val Leu Glu Asn Val Thr Glu Asn
40 45 50

TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAG ATG CAT 193
Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His
55 60

GAG GAT ATA ATC AGT TTA TGG GAT CAA AGT CTA AAG CCA 232
Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys Pro
65 70 75

TGT GTA AAA TTA ACC CCA CTC TGT GTT ACT TTA AAT TGC 271
Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys
80 85 90

ACT AAT TTG GAG AAT GCT AAT AAT ACC GAG AAT GCT AAT 310
Thr Asn Leu Glu Asn Ala Asn Asn Thr Glu Asn Ala Asn
95 100

AAT ACC AAT AAT TAT ACC TTG GGG ATG GAG AGA GGT GAA 349
Asn Thr Asn Asn Tyr Thr Leu Gly Met Glu Arg Gly Glu
105 110 115

AGA AAA AAC TGC TCT TTC AAT ATC ACC ACA AGC TTA AGA 388
Arg Lys Asn Cys Ser Phe Asn Ile Thr Thr Ser Leu Arg
120 125

GAT AAG GGG AAA AAA GAA TAT GCA TTG TTT TAT AAA CTT 427
Asp Lys Gly Lys Lys Glu Tyr Ala Leu Phe Tyr Lys Leu
130 135 140

GAT GTA GTA CAA ATA GAT AAT AGT ACC AAC TAT AGG CTG 466
Asp Val Val Gln Ile Asp Asn Ser Thr Asn Tyr Arg Leu
145 150 155

ATA AGT TGT AAT ACC TCA GTC ATT ACA CAG GCC TGT CCA 505
Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys Pro
160 165

| | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|------|
| | AAG | GTA | TCC | TTT | GAG | CCA | ATT | CCC | ATA | CAT | TAT | TGT | GCC | 544 |
| | Lys | Val | Ser | Phe | Glu | Pro | Ile | Pro | Ile | His | Tyr | Cys | Ala | |
| | 170 | | | | | | 175 | | | | | | 180 | |
| 5 | CCG | GCT | GGT | TTT | GCG | ATT | CTA | AAG | TGT | AAA | GAT | AAG | AAG | 583 |
| | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Lys | Asp | Lys | Lys | |
| | | | | 185 | | | | | 190 | | | | | |
| 10 | TTC | AAT | GGA | ACA | GGA | CCA | TGT | AAA | AAT | GTC | AGG | ACA | GTA | 622 |
| | Phe | Asn | Gly | Thr | Gly | Pro | Cys | Lys | Asn | Val | Arg | Thr | Val | |
| | 195 | | | | | 200 | | | | | 205 | | | |
| 15 | CAA | TGT | ACA | CAT | GGA | ATT | AGA | CCA | GTA | GTA | TCA | ACT | CAA | 661 |
| | Gln | Cys | Thr | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | Gln | |
| | | | 210 | | | | | 215 | | | | | 220 | |
| 20 | CTA | CTG | TTA | AAT | GGC | AGT | CTA | GCA | GAA | GAA | GAG | ATA | GTA | 700 |
| | Leu | Leu | Leu | Asn | Gly | Ser | Leu | Ala | Glu | Glu | Glu | Ile | Val | |
| | | | | | 225 | | | | | 230 | | | | |
| 25 | ATT | AGA | TCT | GAA | AAT | ATC | ACA | GAC | AAT | GCT | AAA | ACC | ATA | 739 |
| | Ile | Arg | Ser | Glu | Asn | Ile | Thr | Asp | Asn | Ala | Lys | Thr | Ile | |
| | | 235 | | | | | 240 | | | | | 245 | | |
| 30 | ATA | GTG | CAG | CTA | AAT | GAA | TCT | ATA | GTG | ATT | AAT | TGT | ACA | 778 |
| | Ile | Val | Gln | Leu | Asn | Glu | Ser | Ile | Val | Ile | Asn | Cys | Thr | |
| | | | | 250 | | | | | 255 | | | | | |
| 35 | AGA | CCC | AAT | AAC | AAC | ACA | AGA | AAA | AGT | ATA | AAT | ATA | GGA | 817 |
| | Arg | Pro | Asn | Asn | Asn | Thr | Arg | Lys | Ser | Ile | Asn | Ile | Gly | |
| | 260 | | | | | 265 | | | | | 270 | | | |
| 40 | CCA | GGG | AGA | GCA | TTC | TAT | ACA | ACA | GGA | GAC | ATA | ATA | GGA | 856 |
| | Pro | Gly | Arg | Ala | Phe | Tyr | Thr | Thr | Gly | Asp | Ile | Ile | Gly | |
| | | | 275 | | | | | 280 | | | | | 285 | |
| 45 | GAT | ATA | AGA | CAA | GCA | CAT | TGT | AAC | CTT | AGT | AAA | ACA | CAA | 895 |
| | Asp | Ile | Arg | Gln | Ala | His | Cys | Asn | Leu | Ser | Lys | Thr | Gln | |
| | | | | | 290 | | | | | 295 | | | | |
| 50 | TGG | GAA | AAA | ACG | TTA | AGA | CAG | ATA | GCT | ATA | AAA | TTA | GAA | 934 |
| | Trp | Glu | Lys | Thr | Leu | Arg | Gln | Ile | Ala | Ile | Lys | Leu | Glu | |
| | 300 | | | | | | 305 | | | | | 310 | | |
| 55 | GAA | AAA | TTT | AAG | AAT | AAA | ACA | ATA | GCC | TTT | AAT | AAA | TCC | 973 |
| | Glu | Lys | Phe | Lys | Asn | Lys | Thr | Ile | Ala | Phe | Asn | Lys | Ser | |
| | | | | 315 | | | | | 320 | | | | | |
| 60 | TCA | GGA | GGG | GAC | CCA | GAA | ATT | GTA | ATG | CAC | AGT | TTT | AAT | 1012 |
| | Ser | Gly | Gly | Asp | Pro | Glu | Ile | Val | Met | His | Ser | Phe | Asn | |
| | 325 | | | | | 330 | | | | | 335 | | | |
| 65 | TGT | GGA | GGG | GGA | TTT | TTC | TAC | TGT | AGT | ACG | AGA | AAA | CTG | 1051 |
| | Cys | Gly | Gly | Gly | Phe | Phe | Tyr | Cys | Ser | Thr | Arg | Lys | Leu | |
| | | | 340 | | | | | 345 | | | | | 350 | |
| 70 | TTT | AAT | AGT | ACC | TGG | AAT | TTA | ACA | CAA | CCG | TTT | AGT | AAT | 1090 |
| | Phe | Asn | Ser | Thr | Trp | Asn | Leu | Thr | Gln | Pro | Phe | Ser | Asn | |
| | | | | | 355 | | | | | 360 | | | | |
| 75 | ACC | GGG | GAT | CGT | ACT | GAA | GAG | TTA | AAT | ATT | ACA | CTC | CCA | 1129 |
| | Thr | Gly | Asp | Arg | Thr | Glu | Glu | Leu | Asn | Ile | Thr | Leu | Pro | |
| | 365 | | | | | 370 | | | | | | 375 | | |

TGC AGA ATA AAA CAA ATC ATA AAC TTG TGG CAG GAA GTA 1168
 Cys Arg Ile Lys Gln Ile Ile Asn Leu Trp Gln Glu Val
 380 385

5 GGC AAA GCA ATG TAT GCC CCT CCC ATC AGA GGA CAA ATT 1207
 Gly Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile
 390 395 400

10 AGA TGT TCA TCA AAT ATT ACA GGG CTA CTA TTA AGG AGA 1246
 Arg Cys Ser Ser Asn Ile Thr Gly Leu Leu Leu Arg Arg
 405 410 415

15 GAT GGT GGA AGT AAC ACC AGT GAC AAC CAG ACT GAG ACC 1285
 Asp Gly Gly Ser Asn Thr Ser Asp Asn Gln Thr Glu Thr
 420 425

20 TTT AGA CCT GGG GGA GGA GAT ATG AGG GAC AAG TGG AGA 1324
 Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Lys Trp Arg
 430 435 440

AGT GAA TTA TAT AAA TAT AAA GTA GTA AGA ATT GAA CCA 1363
 Ser Glu Leu Tyr Lys Tyr Lys Val Val Arg Ile Glu Pro
 445 450

25 TTA GGA GTA GCA CCC ACC CAG GCA AAG AGA AGA GTG CTG 1402
 Leu Gly Val Ala Pro Thr Gln Ala Lys Arg Arg Val Val
 455 460 465

30 CAA AGA GAA AAA AGA GCA GTG GGG ATA GGA GCT ATG TTC 1441
 Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Met Phe
 470 475 480

35 CTT AGG TTC TTA GGA GAT AAA GCT TCT AGA GTC 1474
 Leu Arg Phe Leu Gly Asp Lys Ala Ser Arg Val
 485 490 491

(2) INFORMATION FOR SEQ ID NO:8:
 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 491 amino acids
 (B) TYPE: Amino Acid
 (D) TOPOLOGY: Linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:8:

40 Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe Cys Ala Ser
 1 5 10 15
 Asp Ala Lys Ala Tyr Asp Thr Glu Val His Asn Val Trp Ala Thr
 20 25 30
 50 His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Val Val Leu
 35 40 45
 Glu Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val
 50 55 60
 55 Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu
 65 70 75
 60 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys
 80 85 90
 Thr Asn Leu Glu Asn Ala Asn Asn Thr Glu Asn Ala Asn Asn Thr
 95 100 105

| | | | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|--|
| | Asn | Asn | Tyr | Thr | Leu | Gly | Met | Glu | Arg | Gly | Glu | Arg | Lys | Asn | Cys | |
| | | | | | 110 | | | | | 115 | | | | | 120 | |
| 5 | Ser | Phe | Asn | Ile | Thr | Thr | Ser | Leu | Arg | Asp | Lys | Gly | Lys | Lys | Glu | |
| | | | | | 125 | | | | | 130 | | | | | 135 | |
| | Tyr | Ala | Leu | Phe | Tyr | Lys | Leu | Asp | Val | Val | Gln | Ile | Asp | Asn | Ser | |
| | | | | | 140 | | | | | 145 | | | | | 150 | |
| 10 | Thr | Asn | Tyr | Arg | Leu | Ile | Ser | Cys | Asn | Thr | Ser | Val | Ile | Thr | Gln | |
| | | | | | 155 | | | | | 160 | | | | | 165 | |
| | Ala | Cys | Pro | Lys | Val | Ser | Phe | Glu | Pro | Ile | Pro | Ile | His | Tyr | Cys | |
| 15 | | | | | 170 | | | | | 175 | | | | | 180 | |
| | Ala | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Lys | Asp | Lys | Lys | Phe | |
| | | | | | 185 | | | | | 190 | | | | | 195 | |
| 20 | Asn | Gly | Thr | Gly | Pro | Cys | Lys | Asn | Val | Arg | Thr | Val | Gln | Cys | Thr | |
| | | | | | 200 | | | | | 205 | | | | | 210 | |
| | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | Gln | Leu | Leu | Leu | Asn | Gly | |
| | | | | | 215 | | | | | 220 | | | | | 225 | |
| 25 | Ser | Leu | Ala | Glu | Glu | Glu | Ile | Val | Ile | Arg | Ser | Glu | Asn | Ile | Thr | |
| | | | | | 230 | | | | | 235 | | | | | 240 | |
| | Asp | Asn | Ala | Lys | Thr | Ile | Ile | Val | Gln | Leu | Asn | Glu | Ser | Ile | Val | |
| | | | | | 245 | | | | | 250 | | | | | 255 | |
| 30 | Ile | Asn | Cys | Thr | Arg | Pro | Asn | Asn | Asn | Thr | Arg | Lys | Ser | Ile | Asn | |
| | | | | | 260 | | | | | 265 | | | | | 270 | |
| | Ile | Gly | Pro | Gly | Arg | Ala | Phe | Tyr | Thr | Thr | Gly | Asp | Ile | Ile | Gly | |
| 35 | | | | | 275 | | | | | 280 | | | | | 285 | |
| | Asp | Ile | Arg | Gln | Ala | His | Cys | Asn | Leu | Ser | Lys | Thr | Gln | Trp | Glu | |
| | | | | | 290 | | | | | 295 | | | | | 300 | |
| 40 | Lys | Thr | Leu | Arg | Gln | Ile | Ala | Ile | Lys | Leu | Glu | Glu | Lys | Phe | Lys | |
| | | | | | 305 | | | | | 310 | | | | | 315 | |
| | Asn | Lys | Thr | Ile | Ala | Phe | Asn | Lys | Ser | Ser | Gly | Gly | Asp | Pro | Glu | |
| | | | | | 320 | | | | | 325 | | | | | 330 | |
| 45 | Ile | Val | Met | His | Ser | Phe | Asn | Cys | Gly | Gly | Gly | Phe | Phe | Tyr | Cys | |
| | | | | | 335 | | | | | 340 | | | | | 345 | |
| | Ser | Thr | Arg | Lys | Leu | Phe | Asn | Ser | Thr | Trp | Asn | Leu | Thr | Gln | Pro | |
| 50 | | | | | 350 | | | | | 355 | | | | | 360 | |
| | Phe | Ser | Asn | Thr | Gly | Asp | Arg | Thr | Glu | Glu | Leu | Asn | Ile | Thr | Leu | |
| | | | | | 365 | | | | | 370 | | | | | 375 | |
| 55 | Pro | Cys | Arg | Ile | Lys | Gln | Ile | Ile | Asn | Leu | Trp | Gln | Glu | Val | Gly | |
| | | | | | 380 | | | | | 385 | | | | | 390 | |
| | Lys | Ala | Met | Tyr | Ala | Pro | Pro | Ile | Arg | Gly | Gln | Ile | Arg | Cys | Ser | |
| | | | | | 395 | | | | | 400 | | | | | 405 | |
| 60 | Ser | Asn | Ile | Thr | Gly | Leu | Leu | Leu | Arg | Arg | Asp | Gly | Gly | Ser | Asn | |
| | | | | | 410 | | | | | 415 | | | | | 420 | |
| 65 | Thr | Ser | Asp | Asn | Gln | Thr | Glu | Thr | Phe | Arg | Pro | Gly | Gly | Gly | Asp | |
| | | | | | 425 | | | | | 430 | | | | | 435 | |

Met Arg Asp Lys Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val
 440 445 450
 5 Arg Ile Glu Pro Leu Gly Val Ala Pro Thr Gln Ala Lys Arg Arg
 455 460 465
 Val Val Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Met Phe
 470 475 480
 10 Leu Arg Phe Leu Gly Asp Lys Ala Ser Arg Val
 485 490 491

(2) INFORMATION FOR SEQ ID NO:9:

(i) SEQUENCE CHARACTERISTICS:

- 15 (A) LENGTH: 1512 base pairs
 (B) TYPE: Nucleic Acid
 (C) STRANDEDNESS: Single
 (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:9:

20 CTC GAG GTA CCT GTA TGG AAA GAA GCA ACT ACC ACT 36
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr
 1 5 10
 25 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT AAT ACA GAG 75
 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asn Thr Glu
 15 20 25
 30 AAA CAT AAT GTT TGG GCC ACA CAC GCC TGT GTA CCC ACA 114
 Lys His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr
 30 35
 35 GAT CCC AAC CCA CAA GAA GTA GTA TTG GGA AAT GTG ACA 153
 Asp Pro Asn Pro Gln Glu Val Val Leu Gly Asn Val Thr
 40 45 50
 40 GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA 192
 Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln
 55 60
 45 ATG CAT GAA GAT ATA ATC AGT TTA TGG GAT CAA AGT CTA 231
 Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu
 65 70 75
 50 AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT GTT ACT TTA 270
 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu
 80 85 90
 55 AAT TGC ACT GAT GAT TTA GGG AAT GCT ACT AAT ACC AAT 309
 Asn Cys Thr Asp Asp Leu Gly Asn Ala Thr Asn Thr Asn
 95 100
 60 AGT AGT GCC ACT ACC AAT AGT AGT AGT TGG GAA GAA ATG 348
 Ser Ser Ala Thr Thr Asn Ser Ser Ser Trp Glu Glu Met
 105 110 115
 AAG GGG GAA ATG AAA AGA TGC TCT TTC AAT ATC ACC ACA 387
 Lys Gly Glu Met Lys Arg Cys Ser Phe Asn Ile Thr Thr
 120 125
 60 AGC ATA AGA GAT AAG ATT AAG AAA GAA CAT GCA CTT TTC 426
 Ser Ile Arg Asp Lys Ile Lys Lys Glu His Ala Leu Phe
 130 135 140

| | | |
|----|---|------|
| | TAT AGA CTT GAT GTA GTA CCA ATA GAT AAT GAT AAT ACC | 465 |
| | Tyr Arg Leu Asp Val Val Pro Ile Asp Asn Asp Asn Thr | 155 |
| 5 | ACA TAT AGG TTG ATA AAT TGT AAT ACC TCA GTC ATT ACA | 504 |
| | Thr Tyr Arg Leu Ile Asn Cys Asn Thr Ser Val Ile Thr | 165 |
| 10 | CAG GCC TGT CCA AAG GTA TCA TTT GAG CCA ATT CCC ATA | 543 |
| | Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile | 180 |
| 15 | CAT TTT TGT GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT | 582 |
| | His Phe Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys | 190 |
| 20 | AAT AAT AAG ACG TTC GAG GGA AAA GGA CCA TGT AAA AAT | 621 |
| | Asn Asn Lys Thr Phe Glu Gly Lys Gly Pro Cys Lys Asn | 205 |
| 25 | GTC AGT ACA GTA CAA TGC ACA CAT GGA ATT AGG CCA GTA | 660 |
| | Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val | 220 |
| 30 | GAA GAG GTA ATA ATT AGA TCT GAC AAT ATC ACA GAC AAT | 738 |
| | Glu Glu Val Ile Ile Arg Ser Asp Asn Ile Thr Asp Asn | 245 |
| 35 | ACT AAA ACC ATT ATA GTA CAG CTA AAC GAA TCT GTA GTA | 777 |
| | Thr Lys Thr Ile Ile Val Gln Leu Asn Glu Ser Val Val | 255 |
| 40 | ATT AAT TGT ACA AGA CCC AAC AAC AAT ACA AGA AAA AGT | 816 |
| | Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr Arg Lys Ser | 270 |
| 45 | ATA CAT ATA GGA CCA GGG AGT GCA TTT TTT GCA ACA GGA | 855 |
| | Ile His Ile Gly Pro Gly Ser Ala Phe Phe Ala Thr Gly | 285 |
| 50 | GAA ATA ATA GGA GAT ATA AGA CAA GCA CAC TGT AAC CTT | 894 |
| | Glu Ile Ile Gly Asp Ile Arg Gln Ala His Cys Asn Leu | 295 |
| 55 | AGT AGA ACA CAA TGG AAT AAC ACT TTA GGA AAG ATA GTC | 933 |
| | Ser Arg Thr Gln Trp Asn Asn Thr Leu Gly Lys Ile Val | 310 |
| 60 | ATA AAA TTA AGA GAA CAA TTT AGA AAA CAA TTT GGA GAA | 972 |
| | Ile Lys Leu Arg Glu Gln Phe Arg Lys Gln Phe Gly Glu | 320 |
| | AAA ACA ATA GTC TTT AAT CGA TCC TCA GGA GGG GAC CCG | 1011 |
| | Lys Thr Ile Val Phe Asn Arg Ser Ser Gly Asp Pro | 335 |
| | GAA ATT GCA ATG CAC AGT TTT AAT TGT GGA GGG GAA TTT | 1050 |
| | Glu Ile Ala Met His Ser Phe Asn Cys Gly Gly Glu Phe | 350 |

TTC TAC TGT AAC ACA ACA GCA CTG TTT AAT AGT ACC TGG 1089
 Phe Tyr Cys Asn Thr Thr Ala Leu Phe Asn Ser Thr Trp
 355 360

5 AAT GTT ACT AAA GGG TTG AAT AAC ACT GAA GGA AAT AGC 1128
 Asn Val Thr Lys Gly Leu Asn Asn Thr Glu Gly Asn Ser
 365 370 375

10 ACA GGA GAT GAA AAT ATC ATA CTC CCA TGT AGA ATA AAA 1167
 Thr Gly Asp Glu Asn Ile Ile Leu Pro Cys Arg Ile Lys
 380 385

15 CAA ATT ATA AAC ATG TGG CAG GAA GTA GGA AAA GCA ATG 1206
 Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met
 390 395 400

20 TAT GCC CCT CCC ATC AGT GGA CAA ATT AGA TGT TCA TCA 1245
 Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser
 405 410 415

AAC ATT ACA GGG CTG CTA CTA ACA AGA GAT GGT GGT AGT 1284
 Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Ser
 420 425

25 AAG AAC GAG AGC ATC ACC ACC GAG GTC TTC AGA CCT GGA 1323
 Lys Asn Glu Ser Ile Thr Thr Glu Val Phe Arg Pro Gly
 430 435 440

30 GGA GGA GAT ATG AGG GAC AAT TGG ACA AGT GAA TTA TAT 1362
 Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr
 445 450

35 AAA TAT AAA GTA GTA AAA ATT GAA CCA TTA GGA GTA GCG 1401
 Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala
 455 460 465

40 CCC ACC AAG GCA AAG ACA AGA GTG GTG CAG AGA CAA AAA 1440
 Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys
 470 475 480

AGA GCA GTG GGA ACA ATA GGA GCT ATG TTC CTT GGG TTC 1479
 Arg Ala Val Gly Thr Ile Gly Ala Met Phe Leu Gly Phe
 485 490

45 TTG GGA GCA TAA AGC TTC TAG AGT CGA CCT GCA 1512
 Leu Gly Ala Xaa Ser Phe Xaa Ser Arg Pro Ala
 495 500 504

(2) INFORMATION FOR SEQ ID NO:10:
 50 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 504 amino acids
 (B) TYPE: Amino Acid
 (D) TOPOLOGY: Linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:10:
 55 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe Cys
 1 5 10 15
 60 Ala Ser Asp Ala Lys Ala Tyr Asn Thr Glu Lys His Asn Val Trp
 20 25 30
 Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Val
 35 40 45

| | | | | | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| | Val | Leu | Gly | Asn | Val | Thr | Glu | Asn | Phe | Asn | Met | Trp | Lys | Asn | Asn | 50 | 55 | 60 |
| 5 | Met | Val | Glu | Gln | Met | His | Glu | Asp | Ile | Ile | Ser | Leu | Trp | Asp | Gln | 65 | 70 | 75 |
| | Ser | Leu | Lys | Pro | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Val | Thr | Leu | 80 | 85 | 90 |
| 10 | Asn | Cys | Thr | Asp | Asp | Leu | Gly | Asn | Ala | Thr | Asn | Thr | Asn | Ser | Ser | 95 | 100 | 105 |
| | Ala | Thr | Thr | Asn | Ser | Ser | Ser | Trp | Glu | Glu | Met | Lys | Gly | Glu | Met | 110 | 115 | 120 |
| 15 | Lys | Arg | Cys | Ser | Phe | Asn | Ile | Thr | Thr | Ser | Ile | Arg | Asp | Lys | Ile | 125 | 130 | 135 |
| | Lys | Lys | Glu | His | Ala | Leu | Phe | Tyr | Arg | Leu | Asp | Val | Val | Pro | Ile | 140 | 145 | 150 |
| 20 | Asp | Asn | Asp | Asn | Thr | Thr | Tyr | Arg | Leu | Ile | Asn | Cys | Asn | Thr | Ser | 155 | 160 | 165 |
| 25 | Val | Ile | Thr | Gln | Ala | Cys | Pro | Lys | Val | Ser | Phe | Glu | Pro | Ile | Pro | 170 | 175 | 180 |
| | Ile | His | Phe | Cys | Ala | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Asn | 185 | 190 | 195 |
| 30 | Asn | Lys | Thr | Phe | Glu | Gly | Lys | Gly | Pro | Cys | Lys | Asn | Val | Ser | Thr | 200 | 205 | 210 |
| | Val | Gln | Cys | Thr | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | Gln | Leu | 215 | 220 | 225 |
| 35 | Leu | Leu | Asn | Gly | Ser | Leu | Ala | Glu | Glu | Glu | Val | Ile | Ile | Arg | Ser | 230 | 235 | 240 |
| 40 | Asp | Asn | Ile | Thr | Asp | Asn | Thr | Lys | Thr | Ile | Ile | Val | Gln | Leu | Asn | 245 | 250 | 255 |
| | Glu | Ser | Val | Val | Ile | Asn | Cys | Thr | Arg | Pro | Asn | Asn | Asn | Thr | Arg | 260 | 265 | 270 |
| 45 | Lys | Ser | Ile | His | Ile | Gly | Pro | Gly | Ser | Ala | Phe | Phe | Ala | Thr | Gly | 275 | 280 | 285 |
| | Glu | Ile | Ile | Gly | Asp | Ile | Arg | Gln | Ala | His | Cys | Asn | Leu | Ser | Arg | 290 | 295 | 300 |
| 50 | Thr | Gln | Trp | Asn | Asn | Thr | Leu | Gly | Lys | Ile | Val | Ile | Lys | Leu | Arg | 305 | 310 | 315 |
| 55 | Glu | Gln | Phe | Arg | Lys | Gln | Phe | Gly | Glu | Lys | Thr | Ile | Val | Phe | Asn | 320 | 325 | 330 |
| | Arg | Ser | Ser | Gly | Gly | Asp | Pro | Glu | Ile | Ala | Met | His | Ser | Phe | Asn | 335 | 340 | 345 |
| 60 | Cys | Gly | Gly | Glu | Phe | Phe | Tyr | Cys | Asn | Thr | Thr | Ala | Leu | Phe | Asn | 350 | 355 | 360 |
| 65 | Ser | Thr | Trp | Asn | Val | Thr | Lys | Gly | Leu | Asn | Asn | Thr | Glu | Gly | Asn | 365 | 370 | 375 |

Ser Thr Gly Asp Glu Asn Ile Ile Leu Pro Cys Arg Ile Lys Gln
 380 385 390
 5 Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met Tyr Ala Pro
 395 400 405
 Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser Asn Ile Thr Gly Leu
 410 415 420
 10 Leu Leu Thr Arg Asp Gly Gly Ser Lys Asn Glu Ser Ile Thr Thr
 425 430 435
 Glu Val Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp Arg
 440 445 450
 15 Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly
 455 460 465
 Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys
 470 475 480
 20 Arg Ala Val Gly Thr Ile Gly Ala Met Phe Leu Gly Phe Leu Gly
 485 490 495
 25 Ala Xaa Ser Phe Xaa Ser Arg Pro Ala
 500 504

(2) INFORMATION FOR SEQ ID NO:11:

(i) SEQUENCE CHARACTERISTICS:

- 30 (A) LENGTH: 1501 base pairs
 (B) TYPE: Nucleic Acid
 (C) STRANDEDNESS: Single
 (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:11:

35 CTC GAG GTA CCT GTG TGG AAA GAA GCA ACT ACC ACT 36
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr
 1 5 10
 40 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT AAT ACA GAG 75
 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asn Thr Glu
 15 20 25
 45 AAA CAT AAT GTT TGG GCC ACA CAC GCC TGT GTA CCC ACA 114
 Lys His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr
 30 35
 50 GAT CCC AAC CCA CAA GAA GTA GTA TTG GGA AAT GTG ACA 153
 Asp Pro Asn Pro Gln Glu Val Val Leu Gly Asn Val Thr
 40 45 50
 55 GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA 192
 Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln
 55 60
 60 ATG CAT GAA GAT ATA ATC AGT TTA TGG GAT CAA AGT CTA 231
 Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu
 65 70 75
 AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT GTT ACT TTA 270
 Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu
 80 85 90

AAT TGC ACT GAT GAT TTA GGG AAT GCT ACT AAT ACC AAT 309
 Asn Cys Thr Asp Asp Leu Gly Asn Ala Thr Asn Thr Asn
 95 100

5 AGC AGT GCC ACT ACC AAT AGT AGT AGT TGG GAA GAA ATG 348
 Ser Ser Ala Thr Thr Asn Ser Ser Ser Trp Glu Glu Met
 105 110 115

10 AAG GGG GAA ATG AAA AGG TGC TCT TTC AAT ATC ACC ACA 387
 Lys Gly Glu Met Lys Arg Cys Ser Phe Asn Ile Thr Thr
 120 125

15 AGC ATA AGA GAT AAG ATT AAG AAA GAA CAT GCA CTT TTC 426
 Ser Ile Arg Asp Lys Ile Lys Lys Glu His Ala Leu Phe
 130 135 140

20 TAT AGA CTT GAT GTA GTA CCA ATA GAT AAT GAT AAT ACC 465
 Tyr Arg Leu Asp Val Val Pro Ile Asp Asn Asp Asn Thr
 145 150 155

ACA TAT AGG TTG ATA AAT TGT AAT ACC TCA GTC ATT ACA 504
 Thr Tyr Arg Leu Ile Asn Cys Asn Thr Ser Val Ile Thr
 160 165

25 CAG GCC TGT CCA AAG GTA TCA TTT GAG CCA ATT CCC ATA 543
 Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile
 170 175 180

30 CAT TTT TGT GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT 582
 His Phe Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys
 185 190

35 AAT AAT AAG ACG TTC GAG GGA AAA GGA CCA TGT AAA AAT 621
 Asn Asn Lys Thr Phe Glu Gly Lys Gly Pro Cys Lys Asn
 195 200 205

40 GTC AGT ACA GTA CAA TGC ACA CAT GGA ATT AGG CCA GTA 660
 Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro Val
 210 215 220

GTG TCA ACT CAA CTG CTG TTA AAT GGC AGT CTA GCA GAA 699
 Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu
 225 230

45 GAA GAG GTA ATA ATT AGA TCT GGC AAT ATC ACA GAC AAT 738
 Glu Glu Val Ile Ile Arg Ser Gly Asn Ile Thr Asp Asn
 235 240 245

50 ACT AAA ACC ATT ATA GTA CAG CTA AAC GAA TCT GTA GTA 777
 Thr Lys Thr Ile Ile Val Gln Leu Asn Glu Ser Val Val
 250 255

55 ATT AAT TGT ACA AGA TCC AAC AAC AAT ACA AGA AAA AGT 816
 Ile Asn Cys Thr Arg Ser Asn Asn Asn Thr Arg Lys Ser
 260 265 270

ATA CAT ATA GGA CCA GGG AGT GCA TTT TTT GCA ACA GGA 855
 Ile His Ile Gly Pro Gly Ser Ala Phe Phe Ala Thr Gly
 275 280 285

60 GAA ATA ATA GGA GAT ATA AGA CAA GCA CAC TGT AAC CTT 894
 Glu Ile Ile Gly Asp Ile Arg Gln Ala His Cys Asn Leu
 290 295

AGT AGA ACA CAA TGG AAT AAC ACT TTA GGA AAG ATA GTC 933
 Ser Arg Thr Gln Trp Asn Asn Thr Leu Gly Lys Ile Val
 300 305 310

5 ATA AAA TTA AGA GAA CAA TTT AGA AAA CAA TTT GGA GAA 972
 Ile Lys Leu Arg Glu Gln Phe Arg Lys Gln Phe Gly Glu
 315 320

10 AAA ACA ATA GTC TTT AAT CGA TCC TCA GGA GGG GAC CCG 1011
 Lys Thr Ile Val Phe Asn Arg Ser Ser Gly Gly Asp Pro
 325 330 335

15 GAA ATT GCA ATG CAC AGT TTT AAT TGT GGA GGG GAA TTT 1050
 Glu Ile Ala Met His Ser Phe Asn Cys Gly Gly Glu Phe
 340 345 350

20 TTC TAC TGT AAC ACA ACA GCA CTG TTT AAT AGT ACC TGG 1089
 Phe Tyr Cys Asn Thr Thr Ala Leu Phe Asn Ser Thr Trp
 355 360

AAT GTT ACT AAA GGG TTG AAT AAC ACT GAA GGA AAT AGC 1128
 Asn Val Thr Lys Gly Leu Asn Asn Thr Glu Gly Asn Ser
 365 370 375

25 ACA GGG GAT GAA AAT ATC ATA CTC CCA TGT AGA ATA AAA 1167
 Thr Gly Asp Glu Asn Ile Ile Leu Pro Cys Arg Ile Lys
 380 385

30 CAA ATT ATA AAC ATG TGG CAG GAA GTA GGA AAA GCA ATG 1206
 Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met
 390 395 400

35 TAT GCC CCT CCC ATC AGT GGA CAA ATT AGA TGT TCA TCA 1245
 Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser Ser
 405 410 415

40 AAT ATT ACA GGG CTG CTA CTA ACA AGA GAT GGT GGT AGT 1284
 Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Ser
 420 425

AAG AAC GAG AGC ATC ACC ACC GAG GTC TTC AGA CCT GGA 1323
 Lys Asn Glu Ser Ile Thr Thr Glu Val Phe Arg Pro Gly
 430 435 440

45 GGA GGA GAT ATG AGG GAC AAT TCG AGA AGT GAA TTA TAT 1362
 Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr
 445 450

50 AAA TAT AAA GTA GTA AAA ATT GAA CCA TTA GGA GTA GCG 1401
 Lys Tyr Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala
 455 460 465

55 CCC ACC AAG GCA AAG AGA AGA GTG GTG CAG AGA GAA AAA 1440
 Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys
 470 475 480

60 AGA GCA GTG GGA ACA ATA GGA GCT ATG TTC CTT GGG TTC 1479
 Arg Ala Val Gly Thr Ile Gly Ala Met Phe Leu Gly Phe
 485 490

TTA GGA GCA TAA AGC TTC TAG A 1501
 Leu Gly Ala Xaa Ser Phe Xaa
 495 500

(2) INFORMATION FOR SEQ ID NO:12:

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 500 amino acids

(B) TYPE: Amino Acid

(D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:12:

| | | |
|----|---|-----|
| 5 | Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe Cys | 15 |
| | 1 5 10 | |
| 10 | Ala Ser Asp Ala Lys Ala Tyr Asn Thr Glu Lys His Asn Val Trp | 30 |
| | 20 25 | |
| 15 | Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Val | 45 |
| | 35 40 | |
| | Val Leu Gly Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn | 60 |
| | 50 55 | |
| 20 | Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln | 75 |
| | 65 70 | |
| | Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu | 90 |
| | 80 85 | |
| 25 | Asn Cys Thr Asp Asp Leu Gly Asn Ala Thr Asn Thr Asn Ser Ser | 105 |
| | 95 100 | |
| 30 | Ala Thr Thr Asn Ser Ser Ser Trp Glu Glu Met Lys Gly Glu Met | 120 |
| | 110 115 | |
| | Lys Arg Cys Ser Phe Asn Ile Thr Thr Ser Ile Arg Asp Lys Ile | 135 |
| | 125 130 | |
| 35 | Lys Lys Glu His Ala Leu Phe Tyr Arg Leu Asp Val Val Pro Ile | 150 |
| | 140 145 | |
| | Asp Asn Asp Asn Thr Thr Tyr Arg Leu Ile Asn Cys Asn Thr Ser | 165 |
| | 155 160 | |
| 40 | Val Ile Thr Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro | 180 |
| | 170 175 | |
| 45 | Ile His Phe Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn | 195 |
| | 185 190 | |
| | Asn Lys Thr Phe Glu Gly Lys Gly Pro Cys Lys Asn Val Ser Thr | 210 |
| | 200 205 | |
| 50 | Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr Gln Leu | 225 |
| | 215 220 | |
| | Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val Ile Ile Arg Ser | 240 |
| | 230 235 | |
| 55 | Gly Asn Ile Thr Asp Asn Thr Lys Thr Ile Ile Val Gln Leu Asn | 255 |
| | 245 250 | |
| 60 | Glu Ser Val Val Ile Asn Cys Thr Arg Ser Asn Asn Asn Thr Arg | 270 |
| | 260 265 | |
| | Lys Ser Ile His Ile Gly Pro Gly Ser Ala Phe Phe Ala Thr Gly | 285 |
| | 275 280 | |

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GTA CCC ACA GAC CCT AGT CCA CAA GAA GTA GTT TTG GAA 155
 Val Pro Thr Asp Pro Ser Pro Gln Glu Val Val Leu Glu
 40 45 50

5 AAT GTG ACA GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG 194
 Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn Met
 55 60

10 GTA GAA CAA ATG CAT GAG GAT ATA ATT AGT TTA TGG GAT 233
 Val Glu Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp
 65 70 75

15 CAA AGC TTA AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT 272
 Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys
 80 85 90

20 GTT ACT TTA AAT TGC AGT GAT TAT AGG AAT GCT ACT GAT 311
 Val Thr Leu Asn Cys Ser Asp Tyr Arg Asn Ala Thr Asp
 95 100

25 TAT AAG AAT GCT ACT GAT ACC ACT AGT AGT AAC GAG GGA 350
 Tyr Lys Asn Ala Thr Asp Thr Thr Ser Ser Asn Glu Gly
 105 110 115

30 AAG ATG GAG AGA GGA GAA ATA AAA AAC TGC TCT TTC AAT 389
 Lys Met Glu Arg Gly Glu Ile Lys Asn Cys Ser Phe Asn
 120 125

35 ATT ACC ACA AGC ATA AAA AAT AAG ATG CAG AAA GAA TAT 428
 Ile Thr Thr Ser Ile Lys Asn Lys Met Gln Lys Glu Tyr
 130 135 140

40 GCA CTT TTC TAT AAA CTT GAT ATA GTA CCA ATA GAT AAT 467
 Ala Leu Phe Tyr Lys Leu Asp Ile Val Pro Ile Asp Asn
 145 150 155

45 ACA AGC TAT ACA TTG ATA AGT TGT AAC ACC TCA GTC ATT 506
 Thr Ser Tyr Thr Leu Ile Ser Cys Asn Thr Ser Val Ile
 160 165

50 ACA CAG GCC TGT CCA AAG GTA TCC TTT GAA CCA ACT CCC 545
 Thr Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Thr Pro
 170 175 180

55 ATA CAT TAT TGT GCT CCG GCT GGT TTT GCG ATT CTA AAG 584
 Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys
 185 190

60 TGT AAT GAT AAG AAG TTC AGT GGA AAA GGA GAA TGT AAA 623
 Cys Asn Asp Lys Lys Phe Ser Gly Lys Gly Glu Cys Lys
 195 200 205

65 AAT GTC AGC ACA GTA CAA TGT ACA CAT GGA ATT AGG CCA 662
 Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro
 210 215 220

70 GTA GTA TCA ACT CAA CTG CTG TTA AAT GGC AGT CTA GCA 701
 Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala
 225 230

75 GAA GAA GAG GTG GTA ATT AGA TCT GAC AAT TTC ATA GAC 740
 Glu Glu Glu Val Val Ile Arg Ser Asp Asn Phe Ile Asp
 235 240 245

AAT ACT AAA ACC ATA ATA GTA CAG CTG AAA GAA TCT GTA 779
 Asn Thr Lys Thr Ile Ile Val Gln Leu Lys Glu Ser Val
 250 255

5 GAA ATT AAT TGT ATA AGA CCC AAC AAT AAT ACA AGA AAA 818
 Glu Ile Asn Cys Ile Arg Pro Asn Asn Asn Thr Arg Lys
 260 265 270

10 GGT ATA CAT ATA GGA CCA GGG AGA GCA TGG TAT GCA ACA 857
 Gly Ile His Ile Gly Pro Gly Arg Ala Trp Tyr Ala Thr
 275 280 285

15 GGA GAA ATA GTA GGA GAT ATA AGA AAG CCA TAT TGT AAC 896
 Gly Glu Ile Val Gly Asp Ile Arg Lys Ala Tyr Cys Asn
 290 295

20 ATT AGT AGA ACA AAA TGG AAT AAC ACT TTA ATA CAG ATA 935
 Ile Ser Arg Thr Lys Trp Asn Asn Thr Leu Ile Gln Ile
 300 305 310

GCT AAC AAA TTA AAA GAA AAA TAT AAT ACA ACA ATA AGC 974
 Ala Asn Lys Leu Lys Glu Lys Tyr Asn Thr Thr Ile Ser
 315 320

25 TTT AAT CGA TCC TCA GGA GGG GAC CCA GAA ATT GTA ACG 1013
 Phe Asn Arg Ser Ser Gly Gly Asp Pro Glu Ile Val Thr
 325 330 335

30 CAT AGT TTT AAT TGT GGA GGG GAG TTT TTC TAC TGT GAT 1052
 His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asp
 340 345 350

35 TCA ACA CAA CTG TTT AAT AGT ACT TGG AAT TTA AAT GGT 1091
 Ser Thr Gln Leu Phe Asn Ser Thr Trp Asn Leu Asn Gly
 355 360

40 ACT TGG AAT TTT ACT GCA GGG TCA AAT GAA ACT GAA GGC 1130
 Thr Trp Asn Phe Thr Ala Gly Ser Asn Glu Thr Glu Gly
 365 370 375

AAT ATC ACA CTC CCA TGC AGA ATA AAA CAA ATT ATA AAC 1169
 Asn Ile Thr Leu Pro Cys Arg Ile Lys Gln Ile Ile Asn
 380 385

45 AGG TGG CAG GAA GTA GGG AAA GCA ATG TAT GCC CCT CCC 1208
 Arg Trp Gln Glu Val Gly Lys Ala Met Tyr Ala Pro Pro
 390 395 400

50 ATC AGT GGA CAA ATA AAA TGC TCA TCA AAC ATT ACA GGG 1247
 Ile Ser Gly Gln Ile Lys Cys Ser Ser Asn Ile Thr Gly
 405 410 415

55 ATG ATA TTA ACA AGG GAT GGT GGT AAC GAG AAC AAT AAT 1286
 Met Ile Leu Thr Arg Asp Gly Gly Asn Glu Asn Asn Asn
 420 425

60 GAG AGC AGT ACT ACT GAG ACC TTC AGA CCG GGA GGA GGA 1325
 Glu Ser Ser Thr Thr Glu Thr Phe Arg Pro Gly Gly Gly
 430 435 440

GAT ATG AGG AAC AAT TGG AGA AGT GAA TTA TAT AAA TAT 1364
 Asp Met Arg Asn Asn Trp Arg Ser Glu Leu Tyr Lys Tyr
 445 450

AAA GTA GTA AAA ATT GAA CCA TTA GGA GTA GCA CCC ACC 1403
 Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala Pro Thr
 455 460 465

5 AAG GCA AAG AGA AGA GTG GTG CAG AGA GAA AAA AGA GCA 1442
 Lys Ala Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala
 470 475 480

10 GTG GGA GCG CTA GGA GCT ATG TTC CTT GGG TTC TTA GGA 1481
 Val Gly Ala Leu Gly Ala Met Phe Leu Gly Phe Leu Gly
 485 490

15 GCA TAA ACC TTC TAG ACC GAC TCT AGA GGA TCC 1514
 Ala Xaa Ser Phe Xaa Thr Asp Ser Arg Gly Ser
 495 500 504

(2) INFORMATION FOR SEQ ID NO:14:
 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 504 amino acids
 (B) TYPE: Amino Acid
 (D) TOPOLOGY: Linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:14:

20 Glu Phe Gly Ser Gly Val Pro Val Trp Lys Glu Ala Thr Thr Thr
 1 5 10 15
 Leu Phe Cys Ala Ser Asp Ala Arg Ala Tyr Asp Thr Glu Val His
 20 25 30
 30 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Ser Pro
 35 40 45
 Gln Glu Val Val Leu Glu Asn Val Thr Glu Asn Phe Asn Met Trp
 50 55 60
 35 Lys Asn Asn Met Val Glu Gln Met His Glu Asp Ile Ile Ser Leu
 65 70 75
 40 Trp Asp Gln Ser Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys
 80 85 90
 Val Thr Leu Asn Cys Ser Asp Tyr Arg Asn Ala Thr Asp Tyr Lys
 95 100 105
 45 Asn Ala Thr Asp Thr Thr Ser Ser Asn Glu Gly Lys Met Glu Arg
 110 115 120
 Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Lys
 125 130 135
 50 Asn Lys Met Gln Lys Glu Tyr Ala Leu Phe Tyr Lys Leu Asp Ile
 140 145 150
 55 Val Pro Ile Asp Asn Thr Ser Tyr Thr Leu Ile Ser Cys Asn Thr
 155 160 165
 Ser Val Ile Thr Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Thr
 170 175 180
 60 Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys
 185 190 195
 Asn Asp Lys Lys Phe Ser Gly Lys Gly Glu Cys Lys Asn Val Ser
 200 205 210
 65

| | | | | |
|----|-----------------|---------------------|-------------------------|-----|
| | Thr Val Gln Cys | Thr His Gly Ile Arg | Pro Val Val Ser Thr Gln | |
| | 215 | | 220 | 225 |
| 5 | Leu Leu Leu Asn | Gly Ser Leu Ala Glu | Glu Glu Val Val Ile Arg | |
| | 230 | | 235 | 240 |
| | Ser Asp Asn Phe | Ile Asp Asn Thr Lys | Thr Ile Ile Val Gln Leu | |
| | 245 | | 250 | 255 |
| 10 | Lys Glu Ser Val | Glu Ile Asn Cys Ile | Arg Pro Asn Asn Asn Thr | |
| | 260 | | 265 | 270 |
| | Arg Lys Gly Ile | His Ile Gly Pro Gly | Arg Ala Trp Tyr Ala Thr | |
| | 275 | | 280 | 285 |
| 15 | Gly Glu Ile Val | Gly Asp Ile Arg Lys | Ala Tyr Cys Asn Ile Ser | |
| | 290 | | 295 | 300 |
| | Arg Thr Lys Trp | Asn Asn Thr Leu Ile | Gln Ile Ala Asn Lys Leu | |
| 20 | | 305 | 310 | 315 |
| | Lys Glu Lys Tyr | Asn Thr Thr Ile Ser | Phe Asn Arg Ser Ser Gly | |
| | 320 | | 325 | 330 |
| 25 | Gly Asp Pro Glu | Ile Val Thr His Ser | Phe Asn Cys Gly Gly Glu | |
| | 335 | | 340 | 345 |
| | Phe Phe Tyr Cys | Asp Ser Thr Gln Leu | Phe Asn Ser Thr Trp Asn | |
| | 350 | | 355 | 360 |
| 30 | Leu Asn Gly Thr | Trp Asn Phe Thr Ala | Gly Ser Asn Glu Thr Glu | |
| | 365 | | 370 | 375 |
| | Gly Asn Ile Thr | Leu Pro Cys Arg Ile | Lys Gln Ile Ile Asn Arg | |
| 35 | | 380 | 385 | 390 |
| | Trp Gln Glu Val | Gly Lys Ala Met Tyr | Ala Pro Pro Ile Ser Gly | |
| | 395 | | 400 | 405 |
| 40 | Gln Ile Lys Cys | Ser Ser Asn Ile Thr | Gly Met Ile Leu Thr Arg | |
| | 410 | | 415 | 420 |
| | Asp Gly Gly Asn | Glu Asn Asn Asn Glu | Ser Ser Thr Thr Glu Thr | |
| | 425 | | 430 | 435 |
| 45 | Phe Arg Pro Gly | Gly Gly Asp Met Arg | Asn Asn Trp Arg Ser Glu | |
| | 440 | | 445 | 450 |
| | Leu Tyr Lys Tyr | Lys Val Val Lys Ile | Glu Pro Leu Gly Val Ala | |
| 50 | | 455 | 460 | 465 |
| | Pro Thr Lys Ala | Lys Arg Arg Val Val | Gln Arg Glu Lys Arg Ala | |
| | 470 | | 475 | 480 |
| 55 | Val Gly Ala Leu | Gly Ala Met Phe Leu | Gly Phe Leu Gly Ala Xaa | |
| | 485 | | 490 | 495 |
| | Ser Phe Xaa Thr | Asp Ser Arg Gly Ser | | |
| | 500 | | 504 | |

(2) INFORMATION FOR SEQ ID NO:15:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1408 base pairs

(B) TYPE: Nucleic Acid

(C) STRANDEDNESS: Single

(D) TOPOLOGY: Linear
 (x1) SEQUENCE DESCRIPTION: SEQ ID NO:15:

| | | | | | | | | | | | | | | |
|----|---|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| 5 | G | GTA | CCT | GTG | TGG | AAG | GAA | GCA | ACC | ACC | ACT | CTA | TTC | 37 |
| | | Val | Pro | Val | Trp | Lys | Glu | Ala | Thr | Thr | Thr | Leu | Phe | |
| | | 1 | | | | 5 | | | | | | 10 | | |
| 10 | | TGT | GCA | TCA | GAT | GCT | AGA | GCA | TAT | GAC | ACA | GAG | GTA | 76 |
| | | Cys | Ala | Ser | Asp | Ala | Arg | Ala | Tyr | Asp | Thr | Glu | Val | His |
| | | | 15 | | | | 20 | | | | | | 25 | |
| 15 | | AAT | GTT | TGG | GCC | ACA | CAT | GCC | TGT | GTA | CCC | ACA | GAC | 115 |
| | | Asn | Val | Trp | Ala | Thr | His | Ala | Cys | Val | Pro | Thr | Asp | Pro |
| | | | | | 30 | | | | | | 35 | | | |
| 20 | | AGT | CCA | CAA | GAA | GTA | TTT | TTG | GGA | AAT | GTG | ACA | GAA | 154 |
| | | Ser | Pro | Gln | Glu | Val | Phe | Leu | Gly | Asn | Val | Thr | Glu | Asn |
| | | | 40 | | | | 45 | | | | | | 50 | |
| 25 | | TTT | AAT | ATG | TGG | AAA | AAT | AAC | ATG | GTA | GAA | CAA | ATG | 193 |
| | | Phe | Asn | Met | Trp | Lys | Asn | Asn | Met | Val | Glu | Gln | Met | Tyr |
| | | | | | 55 | | | | | 60 | | | | |
| 30 | | GAG | GAT | ATA | ATT | AGT | TTA | TGG | GAT | CAA | AGC | TTA | AAG | 232 |
| | | Glu | Asp | Ile | Ile | Ser | Leu | Trp | Asp | Gln | Ser | Leu | Lys | Pro |
| | | | 65 | | | | 70 | | | | | 75 | | |
| 35 | | TGT | GTA | AAA | TTA | ACC | CCA | CTC | TGT | GTT | ACT | TTA | AAT | 271 |
| | | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Val | Thr | Leu | Asn | Cys |
| | | | | 80 | | | | | 85 | | | | 90 | |
| 40 | | AGT | GAT | TAT | AGG | AAT | GCT | ACT | GAT | TAT | AAG | AAT | GCT | 310 |
| | | Ser | Asp | Tyr | Arg | Asn | Ala | Thr | Asp | Tyr | Lys | Asn | Ala | Thr |
| | | | | | 95 | | | | | | 100 | | | |
| 45 | | GAT | ACC | ACT | AGT | AGT | AAC | GAG | GGA | AAG | ATG | GAG | AGA | 349 |
| | | Asp | Thr | Thr | Ser | Ser | Asn | Glu | Gly | Lys | Met | Glu | Arg | Gly |
| | | | 105 | | | | 110 | | | | | | 115 | |
| 50 | | GAA | ATA | AAA | AAC | TGC | TCT | TTC | AAT | ATC | ACC | ACA | AGC | 388 |
| | | Glu | Ile | Lys | Asn | Cys | Ser | Phe | Asn | Ile | Thr | Thr | Ser | Ile |
| | | | | | 120 | | | | | 125 | | | | |
| 55 | | AAA | AAT | AAG | ATG | CAG | AAA | GAA | TAT | GCA | CTT | TTC | TAT | 427 |
| | | Lys | Asn | Lys | Met | Gln | Lys | Glu | Tyr | Ala | Leu | Phe | Tyr | Lys |
| | | | 130 | | | | 135 | | | | | 140 | | |
| 60 | | CTT | AAT | ATA | GTA | CCA | ATA | GAT | AAT | ACA | AGC | TAT | ACA | 466 |
| | | Leu | Asn | Ile | Val | Pro | Ile | Asp | Asn | Thr | Ser | Tyr | Thr | Leu |
| | | | | 145 | | | | | 150 | | | | 155 | |
| 65 | | ATA | AGT | TGT | AAC | ACC | TCA | GTC | ATT | ACA | CAG | GCC | TGT | 505 |
| | | Ile | Ser | Cys | Asn | Thr | Ser | Val | Ile | Thr | Gln | Ala | Cys | Pro |
| | | | | | 160 | | | | | | 165 | | | |
| 70 | | AAG | GTA | TCC | TTT | GAA | CCA | ATT | CCC | ATA | CAT | TAT | TGT | 544 |
| | | Lys | Val | Ser | Phe | Glu | Pro | Ile | Pro | Ile | His | Tyr | Cys | Ala |
| | | | 170 | | | | 175 | | | | | | 180 | |
| 75 | | CCG | GCT | GGT | TTT | GCG | ATT | CTA | AAG | TGT | AAT | GAT | AAG | 583 |
| | | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Asn | Asp | Lys | Lys |
| | | | | | 185 | | | | | 190 | | | | |

| | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|------|
| | TTC | AGT | GGA | AAA | GGA | GAA | TGT | AAA | AAT | CTC | AGC | ACA | GTA | 622 |
| | Phe | Ser | Gly | Lys | Gly | Glu | Cys | Lys | Asn | Val | Ser | Thr | Val | |
| | 195 | | | | | 200 | | | | | 205 | | | |
| 5 | CAA | TGT | ACA | CAT | GGA | ATT | AGG | CCA | GTA | GTA | TCA | ACT | CAA | 661 |
| | Gln | Cys | Thr | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | Gln | |
| | | | 210 | | | | | 215 | | | | | 220 | |
| 10 | CTG | CTG | TTA | AAT | GGC | AGT | CTA | GCA | GAA | GAA | GAG | GTG | GTA | 700 |
| | Leu | Leu | Leu | Asn | Gly | Ser | Leu | Ala | Glu | Glu | Glu | Val | Val | |
| | | | | | 225 | | | | | 230 | | | | |
| 15 | ATT | AGA | TCT | GAC | AAT | TTC | ACA | GAC | AAT | ACT | AAA | ACC | ATA | 739 |
| | Ile | Arg | Ser | Asp | Asn | Phe | Thr | Asp | Asn | Thr | Lys | Thr | Ile | |
| | | 235 | | | | | 240 | | | | | 245 | | |
| 20 | ATA | GTA | CAG | CTG | AAA | GAA | TCT | GTA | GAA | ATT | AAT | TGT | ATA | 778 |
| | Ile | Val | Gln | Leu | Lys | Glu | Ser | Val | Glu | Ile | Asn | Cys | Ile | |
| | | | | 250 | | | | | 255 | | | | | |
| | AGA | CCC | AAC | AAT | AAT | ACA | AGA | AAA | GGT | ATA | CAT | ATA | GGA | 817 |
| | Arg | Pro | Asn | Asn | Asn | Thr | Arg | Lys | Gly | Ile | His | Ile | Gly | |
| | 260 | | | | | 265 | | | | | 270 | | | |
| 25 | CCA | GGG | AGA | GCA | TGG | TAT | GCA | ACA | GGA | GAA | ATA | GTA | GGA | 856 |
| | Pro | Gly | Arg | Ala | Trp | Tyr | Ala | Thr | Gly | Glu | Ile | Val | Gly | |
| | | | 275 | | | | | 280 | | | | | 285 | |
| 30 | GAT | ATA | AGA | CAG | GCA | TAT | TGT | AAC | ATT | AGT | AGA | ACA | AAA | 895 |
| | Asp | Ile | Arg | Gln | Ala | Tyr | Cys | Asn | Ile | Ser | Arg | Thr | Lys | |
| | | | | | 290 | | | | | 295 | | | | |
| 35 | TGG | AAT | AAC | ACT | TTA | ATA | CAG | ATA | GCT | AAC | AAA | TTA | AAA | 934 |
| | Trp | Asn | Asn | Thr | Leu | Ile | Gln | Ile | Ala | Asn | Lys | Leu | Lys | |
| | | 300 | | | | | 305 | | | | | 310 | | |
| 40 | GAA | AAA | TAT | AAT | ACA | ACA | ATA | AGC | TTT | AAT | CGA | TCC | TCA | 973 |
| | Glu | Lys | Tyr | Asn | Thr | Thr | Ile | Ser | Phe | Asn | Arg | Ser | Ser | |
| | | | | 315 | | | | | 320 | | | | | |
| | GGA | GGG | GAC | CCA | GAA | ATT | GTA | ACC | CAT | AGT | TTT | AAT | TGT | 1012 |
| | Gly | Gly | Asp | Pro | Glu | Ile | Val | Thr | His | Ser | Phe | Asn | Cys | |
| | 325 | | | | | 330 | | | | | 335 | | | |
| 45 | GGA | GGG | GAA | TTT | TTC | TAC | TGT | AAT | TCA | ACA | CAA | CTG | TTT | 1051 |
| | Gly | Gly | Glu | Phe | Phe | Tyr | Cys | Asn | Ser | Thr | Gln | Leu | Phe | |
| | | | 340 | | | | | 345 | | | | | 350 | |
| 50 | AAT | AGT | ACT | TGG | AAT | TTA | AAT | GGT | ACT | TGG | AAT | TTT | ACT | 1090 |
| | Asn | Ser | Thr | Trp | Asn | Leu | Asn | Gly | Thr | Trp | Asn | Phe | Thr | |
| | | | | | 355 | | | | | 360 | | | | |
| 55 | GCA | GGG | TCA | AAT | GAA | ACT | GAA | GGC | AAT | ATC | ACA | CTC | CCA | 1129 |
| | Ala | Gly | Ser | Asn | Glu | Thr | Glu | Gly | Asn | Ile | Thr | Leu | Pro | |
| | | 365 | | | | | | 370 | | | | 375 | | |
| 60 | TGC | AGA | ATA | AAA | CAA | ATT | ATA | AAC | ACG | TGG | CAG | GAA | GTA | 1168 |
| | Cys | Arg | Ile | Lys | Gln | Ile | Ile | Asn | Arg | Trp | Gln | Glu | Val | |
| | | | | 380 | | | | | 385 | | | | | |
| | GGA | AAA | GCA | ATG | TAT | GCC | CCT | CCC | ATC | AGT | GGA | CAA | ATA | 1207 |
| | Gly | Lys | Ala | Met | Tyr | Ala | Pro | Pro | Ile | Ser | Gly | Gln | Ile | |
| | 390 | | | | | 395 | | | | | 400 | | | |

AGA TGC TCA TCA AAC ATT ACA GGG ATG ATA TTA ACA AGG 1246
 Arg Cys Ser Ser Asn Ile Thr Gly Met Ile Leu Thr Arg
 405 410 415

5 GAT GGT GGT AAC GAG AAC AAT AAT GAG AGC AGT ACT ACT 1285
 Asp Gly Gly Asn Glu Asn Asn Asn Glu Ser Ser Thr Thr
 420 425

10 GAG ACC TTC AGA CCG GGA GGA GGA GAT ATG AGG AAC AAT 1324
 Glu Thr Phe Arg Pro Gly Gly Gly Asp Met Arg Asn Asn
 430 435 440

15 TGG AGA AGT GAA TTA TAT AAA TAT AAA GTA GTA AAA ATT 1363
 Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile
 445 450

GAG CCA TTA GGA GTA GCA CCC ACC GAC TCT AGA GGA TCC 1402
 Glu Pro Leu Gly Val Ala Pro Thr Asp Ser Arg Gly Ser
 455 460 465

20 TCT AGA 1408
 Ser Arg
 469

25 (2) INFORMATION FOR SEQ ID NO:16:
 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 469 amino acids
 (B) TYPE: Amino Acid
 (D) TOPOLOGY: Linear

30 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:16:

Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe Cys Ala Ser
 1 5 10 15

35 Asp Ala Arg Ala Tyr Asp Thr Glu Val His Asn Val Trp Ala Thr
 20 25 30

His Ala Cys Val Pro Thr Asp Pro Ser Pro Gln Glu Val Phe Leu
 35 40 45

40 Gly Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val
 50 55 60

45 Glu Gln Met Tyr Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu
 65 70 75

Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys
 80 85 90

50 Ser Asp Tyr Arg Asn Ala Thr Asp Tyr Lys Asn Ala Thr Asp Thr
 95 100 105

Thr Ser Ser Asn Glu Gly Lys Met Glu Arg Gly Glu Ile Lys Asn
 110 115 120

55 Cys Ser Phe Asn Ile Thr Thr Ser Ile Lys Asn Lys Met Gln Lys
 125 130 135

Glu Tyr Ala Leu Phe Tyr Lys Leu Asn Ile Val Pro Ile Asp Asn
 140 145 150

60 Thr Ser Tyr Thr Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln
 155 160 165

| | | | | |
|----|---|-----|-----|-----|
| | Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys | 170 | 175 | 180 |
| 5 | Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asp Lys Lys Phe | 185 | 190 | 195 |
| | Ser Gly Lys Gly Glu Cys Lys Asn Val Ser Thr Val Gln Cys Thr | 200 | 205 | 210 |
| 10 | His Gly Ile Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly | 215 | 220 | 225 |
| | Ser Leu Ala Glu Glu Glu Val Val Ile Arg Ser Asp Asn Phe Thr | 230 | 235 | 240 |
| 15 | Asp Asn Thr Lys Thr Ile Ile Val Gln Leu Lys Glu Ser Val Glu | 245 | 250 | 255 |
| | Ile Asn Cys Ile Arg Pro Asn Asn Asn Thr Arg Lys Gly Ile His | 260 | 265 | 270 |
| 20 | Ile Gly Pro Gly Arg Ala Trp Tyr Ala Thr Gly Glu Ile Val Gly | 275 | 280 | 285 |
| | Asp Ile Arg Gln Ala Tyr Cys Asn Ile Ser Arg Thr Lys Trp Asn | 290 | 295 | 300 |
| | Asn Thr Leu Ile Gln Ile Ala Asn Lys Leu Lys Glu Lys Tyr Asn | 305 | 310 | 315 |
| 30 | Thr Thr Ile Ser Phe Asn Arg Ser Ser Gly Gly Asp Pro Glu Ile | 320 | 325 | 330 |
| | Val Thr His Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn | 335 | 340 | 345 |
| 35 | Ser Thr Gln Leu Phe Asn Ser Thr Trp Asn Leu Asn Gly Thr Trp | 350 | 355 | 360 |
| | Asn Phe Thr Ala Gly Ser Asn Glu Thr Glu Gly Asn Ile Thr Leu | 365 | 370 | 375 |
| 40 | Pro Cys Arg Ile Lys Gln Ile Ile Asn Arg Trp Gln Glu Val Gly | 380 | 385 | 390 |
| 45 | Lys Ala Met Tyr Ala Pro Pro Ile Ser Gly Gln Ile Arg Cys Ser | 395 | 400 | 405 |
| | Ser Asn Ile Thr Gly Met Ile Leu Thr Arg Asp Gly Gly Asn Glu | 410 | 415 | 420 |
| 50 | Asn Asn Asn Glu Ser Ser Thr Thr Glu Thr Phe Arg Pro Gly Gly | 425 | 430 | 435 |
| | Gly Asp Met Arg Asn Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys | 440 | 445 | 450 |
| 55 | Val Val Lys Ile Glu Pro Leu Gly Val Ala Pro Thr Asp Ser Arg | 455 | 460 | 465 |
| 60 | Gly Ser Ser Arg | 469 | | |

(2) INFORMATION FOR SEQ ID NO:17:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1499 base pairs

(B) TYPE: Nucleic Acid

(C) STRANDEDNESS: Single

(D) TOPOLOGY: Linear

5 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:17:

GAG GTA CCT GTG TGG AAA GAA GCA ACC ACT ACT CTA 36
 Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu
 1 5 10

10 TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GGG GTG 75
 Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Gly Val
 15 20 25

15 CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA GAC 114
 His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp
 30 35

20 CCC AAC CCA CAA GAA ATA GAA TTG GTA AAT GTG ACA GAA 153
 Pro Asn Pro Gln Glu Ile Glu Leu Val Asn Val Thr Glu
 40 45 50

25 GAT TTT AAC ATG TGG AAA AAT AAA ATG GTA GAC CAG ATG 192
 Asp Phe Asn Met Trp Lys Asn Lys Met Val Asp Gln Met
 55 60

30 CAT GAG GAT ATA ATC AGT TTA TGG GAT GAA AGC CTA AAG 231
 His Glu Asp Ile Ile Ser Leu Trp Asp Glu Ser Leu Lys
 65 70 75

35 CCA TGT GTA AAG TTA ACC CCA CTT TGT GTT ACT CTA AAC 270
 Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn
 80 85 90

40 TGC AGT GAT GTG AAC AAT TCC ACA AAT CCT AAT GAT ACT 309
 Cys Ser Asp Val Asn Asn Ser Thr Asn Pro Asn Asp Thr
 95 100

45 AAT ACT AAT TCC ACT AAT ACT ACT TCC TCT ACT CCT ACG 348
 Asn Thr Asn Ser Thr Asn Thr Thr Ser Ser Thr Pro Thr
 105 110 115

50 GCC ACT ACT AGT AGC GAG GAA AAG ATG GAG AAG GGA GAA 387
 Ala Thr Thr Ser Ser Glu Glu Lys Met Glu Lys Gly Glu
 120 125

55 ATA AAA AAC TGC TCT TTC AAT ATC ACC ACA CAC ATG AAA 426
 Ile Lys Asn Cys Ser Phe Asn Ile Thr Thr His Met Lys
 130 135 140

60 GAT AAG GCA CAG AAA GAA TAT GCA CTT TTT TAT AAA CTT 465
 Asp Lys Ala Gln Lys Glu Tyr Ala Leu Phe Tyr Lys Leu
 145 150 155

65 GAT ATA GTA CCA ATA GAT GAT AAT AAT GCC AGC TAT AGG 504
 Asp Ile Val Pro Ile Asp Asp Asn Asn Ala Ser Tyr Arg
 160 165

70 TTG ATA AGT TGT AAT ACC TCA GAC ATT ACA CAG GCC TGT 543
 Leu Ile Ser Cys Asn Thr Ser Asp Ile Thr Gln Ala Cys
 170 175 180

75 CCA AAG GTG ACC TTT GAG CCA ATT CCC ATA CAT TAT TGT 582
 Pro Lys Val Thr Phe Glu Pro Ile Pro Ile His Tyr Cys
 185 190

GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT AAA GAT AAG 621
 Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Lys Asp Lys
 195 200 205

5 AAG TTC AAT GGA ACA GGA CCA TGT TCA AAG GTC AGC ACA 660
 Lys Phe Asn Gly Thr Gly Pro Cys Ser Lys Val Ser Thr
 210 215 220

10 GTA CAA TGT ACA CAT GGA ATT AGG CCA GTA GTA TCA ACT 699
 Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr
 225 230

15 CAA CTG TTG TTA AAT GGC AGT CTT GCA GAA GAA GAA GTA 738
 Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val
 235 240 245

20 GTA ATT AGA TCT GTC AAT TTC ACA GAC AAT GCT AAA ATC 777
 Val Ile Arg Ser Val Asn Phe Thr Asp Asn Ala Lys Ile
 250 255

25 ATA ATA GTA CAG CTG AAA GAA CCT GTA GCA ATT AAT TGT 816
 Ile Ile Val Gln Leu Lys Glu Pro Val Ala Ile Asn Cys
 260 265 270

30 ACA AGA CCC AAC AAC AAT ACA AGA AAA GGT ATA CAT CTA 855
 Thr Arg Pro Asn Asn Asn Thr Arg Lys Gly Ile His Leu
 275 280 285

35 GGA CCA GGG AGC ACA TTT TAT ACA ACA GGA GAA ATA ATA 894
 Gly Pro Gly Ser Thr Phe Tyr Thr Thr Gly Glu Ile Ile
 290 295

40 GGA GAC ATA AGA AAA GCA TAT TGC AAG ATT AGT AAA GAA 933
 Gly Asp Ile Arg Lys Ala Tyr Cys Lys Ile Ser Lys Glu
 300 305 310

45 AAA TGG AAT AAC ACT TTA AGA CAG GTA GTT AAA AAA TTA 972
 Lys Trp Asn Asn Thr Leu Arg Gln Val Val Lys Lys Leu
 315 320

50 AGA GAA CAA TTT GGG AAT AAA ACA ATA ATT TTT AAT CGA 1011
 Arg Glu Gln Phe Gly Asn Lys Thr Ile Ile Phe Asn Arg
 325 330 335

55 TCC TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC AGT TTT 1050
 Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Ser Phe
 340 345 350

60 AAC TGT GGA GGG GAG TTT TTC TAC TGT AAT ACA ACA CAA 1089
 Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr Gln
 355 360

65 CTG TTT AAT AGT ACT TGG AAT AAT ACT GAA GGG ACA AAT 1128
 Leu Phe Asn Ser Thr Trp Asn Asn Thr Glu Gly Thr Asn
 365 370 375

70 AGC ACT GAA GGA AAT AGC ACA ATC ACA CTC CCA TGC AGA 1167
 Ser Thr Glu Gly Asn Ser Thr Ile Thr Leu Pro Cys Arg
 380 385

75 ATA AAA CAA ATT ATA AAT ATG TGG CAG GAA GTA GGA AAA 1206
 Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 390 395 400

GCA ACG TAT GCC CCT CCC ATC AGA GGA CGA ATT AGA TGC 1245
 Ala Thr Tyr Ala Pro Pro Ile Arg Gly Arg Ile Arg Cys
 405 410 415

5 ATA TCA AAT ATT ACA GGA CTG CTA TTA ACA AGA GAT GGT 1284
 Ile Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly
 420 425

10 GGT AGG AAT GTC ACA AAC AAT ACC GAA ACC TTC AGA CCT 1323
 Gly Arg Asn Val Thr Asn Asn Thr Glu Thr Phe Arg Pro
 430 435 440

15 GGA GGA GGA GAC ATG AGG GAC AAT TGG AGA AGT GAA TTA 1362
 Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu
 445 450

20 TAT AAA TAT AAA GTA GTA AAA GTT GAA CCA TTA GGA ATA 1401
 Tyr Lys Tyr Lys Val Val Lys Val Glu Pro Leu Gly Ile
 455 460 465

GCA CCC ACC AAG GCA AAG AGA AGA GTG GTG CAC AGA GAC 1440
 Ala Pro Thr Lys Ala Lys Arg Arg Val Val His Arg Asp
 470 475 480

25 AAA AGA GCA GCA CTA GGA GCC TTG TTC CTT GGG TTC TTA 1479
 Lys Arg Ala Ala Leu Gly Ala Leu Phe Leu Gly Phe Leu
 485 490

30 GGA GCA TAA AAG CTT CTA GA 1499
 Gly Ala Xaa Lys Leu Leu
 495 499

(2) INFORMATION FOR SEQ ID NO:18:
 (i) SEQUENCE CHARACTERISTICS:
 35 (A) LENGTH: 499 amino acids
 (B) TYPE: Amino Acid
 (D) TOPOLOGY: Linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:18:

40 Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe Cys Ala
 1 5 10 15

Ser Asp Ala Lys Ala Tyr Asp Thr Gly Val His Asn Val Trp Ala
 20 25 30

45 Thr His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Ile Glu
 35 40 45

50 Leu Val Asn Val Thr Glu Asp Phe Asn Met Trp Lys Asn Lys Met
 50 55 60

Val Asp Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp Glu Ser
 65 70 75

55 Leu Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn
 80 85 90

Cys Ser Asp Val Asn Asn Ser Thr Asn Pro Asn Asp Thr Asn Thr
 95 100 105

60 Asn Ser Thr Asn Thr Thr Ser Ser Thr Pro Thr Ala Thr Thr Ser
 110 115 120

65 Ser Glu Glu Lys Met Glu Lys Gly Glu Ile Lys Asn Cys Ser Phe
 125 130 135

| | | | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|--|
| | Asn | Ile | Thr | Thr | His | Met | Lys | Asp | Lys | Ala | Gln | Lys | Glu | Tyr | Ala | |
| | | | | | 140 | | | | | 145 | | | | | 150 | |
| 5 | Leu | Phe | Tyr | Lys | Leu | Asp | Ile | Val | Pro | Ile | Asp | Asp | Asn | Asn | Ala | |
| | | | | | 155 | | | | | 160 | | | | | 165 | |
| | Ser | Tyr | Arg | Leu | Ile | Ser | Cys | Asn | Thr | Ser | Asp | Ile | Thr | Gln | Ala | |
| | | | | | 170 | | | | | 175 | | | | | 180 | |
| 10 | Cys | Pro | Lys | Val | Thr | Phe | Glu | Pro | Ile | Pro | Ile | His | Tyr | Cys | Ala | |
| | | | | | 185 | | | | | 190 | | | | | 195 | |
| | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Lys | Asp | Lys | Lys | Phe | Asn | |
| 15 | | | | | 200 | | | | | 205 | | | | | 210 | |
| | Gly | Thr | Gly | Pro | Cys | Ser | Lys | Val | Ser | Thr | Val | Gln | Cys | Thr | His | |
| | | | | | 215 | | | | | 220 | | | | | 225 | |
| 20 | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | Gln | Leu | Leu | Leu | Asn | Gly | Ser | |
| | | | | | 230 | | | | | 235 | | | | | 240 | |
| | Leu | Ala | Glu | Glu | Glu | Val | Val | Ile | Arg | Ser | Val | Asn | Phe | Thr | Asp | |
| | | | | | 245 | | | | | 250 | | | | | 255 | |
| 25 | Asn | Ala | Lys | Ile | Ile | Ile | Val | Gln | Leu | Lys | Glu | Pro | Val | Ala | Ile | |
| | | | | | 260 | | | | | 265 | | | | | 270 | |
| | Asn | Cys | Thr | Arg | Pro | Asn | Asn | Asn | Thr | Arg | Lys | Gly | Ile | His | Leu | |
| 30 | | | | | 275 | | | | | 280 | | | | | 285 | |
| | Gly | Pro | Gly | Ser | Thr | Phe | Tyr | Thr | Thr | Gly | Glu | Ile | Ile | Gly | Asp | |
| | | | | | 290 | | | | | 295 | | | | | 300 | |
| 35 | Ile | Arg | Lys | Ala | Tyr | Cys | Lys | Ile | Ser | Lys | Glu | Lys | Trp | Asn | Asn | |
| | | | | | 305 | | | | | 310 | | | | | 315 | |
| | Thr | Leu | Arg | Gln | Val | Val | Lys | Lys | Leu | Arg | Glu | Gln | Phe | Gly | Asn | |
| | | | | | 320 | | | | | 325 | | | | | 330 | |
| 40 | Lys | Thr | Ile | Ile | Phe | Asn | Arg | Ser | Ser | Gly | Gly | Asp | Pro | Glu | Ile | |
| | | | | | 335 | | | | | 340 | | | | | 345 | |
| | Val | Met | His | Ser | Phe | Asn | Cys | Gly | Gly | Glu | Phe | Phe | Tyr | Cys | Asn | |
| 45 | | | | | 350 | | | | | 355 | | | | | 360 | |
| | Thr | Thr | Gln | Leu | Phe | Asn | Ser | Thr | Trp | Asn | Asn | Thr | Glu | Gly | Thr | |
| | | | | | 365 | | | | | 370 | | | | | 375 | |
| 50 | Asn | Ser | Thr | Glu | Gly | Asn | Ser | Thr | Ile | Thr | Leu | Pro | Cys | Arg | Ile | |
| | | | | | 380 | | | | | 385 | | | | | 390 | |
| | Lys | Gln | Ile | Ile | Asn | Met | Trp | Gln | Glu | Val | Gly | Lys | Ala | Thr | Tyr | |
| | | | | | 395 | | | | | 400 | | | | | 405 | |
| 55 | Ala | Pro | Pro | Ile | Arg | Gly | Arg | Ile | Arg | Cys | Ile | Ser | Asn | Ile | Thr | |
| | | | | | 410 | | | | | 415 | | | | | 420 | |
| | Gly | Leu | Leu | Leu | Thr | Arg | Asp | Gly | Gly | Arg | Asn | Val | Thr | Asn | Asn | |
| 60 | | | | | 425 | | | | | 430 | | | | | 435 | |
| | Thr | Glu | Thr | Phe | Arg | Pro | Gly | Gly | Gly | Asp | Met | Arg | Asp | Asn | Trp | |
| | | | | | 440 | | | | | 445 | | | | | 450 | |
| 65 | Arg | Ser | Glu | Leu | Tyr | Lys | Tyr | Lys | Val | Val | Lys | Val | Glu | Pro | Leu | |
| | | | | | 455 | | | | | 460 | | | | | 465 | |

Gly Ile Ala Pro Thr Lys Ala Lys Arg Arg Val Val His Arg Asp
 470 475 480
 5 Lys Arg Ala Ala Leu Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala
 485 490 495
 Xaa Lys Leu Leu
 499

10 (2) INFORMATION FOR SEQ ID NO:19:
 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1499 base pairs
 (B) TYPE: Nucleic Acid
 (C) STRANDEDNESS: Single
 15 (D) TOPOLOGY: Linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:19:

GAG GTA CCT GTA TGG AAA GAA GCA ACC ACT ACT CTA 36
 Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu
 1 5 10
 20 TTT TGT GCA TCA GAT GCT AAA GCA TAT GAC ACA GAG GTG 75
 Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val
 15 20 25
 25 CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA GAC 114
 His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp
 30 35
 30 CCC AAC CCA CAA GAA ATA GAA TTG GTA AAT GTG ACA GAA 153
 Pro Asn Pro Gln Glu Ile Glu Leu Val Asn Val Thr Glu
 40 45 50
 35 GAT TTT AAC ATG TGG AAA AAT AAA ATG GTA GAC CAG ATG 192
 Asp Phe Asn Met Trp Lys Asn Lys Met Val Asp Gln Met
 55 60
 40 CAT GAG GAT ATA ATC AGT TTA TGG GAT GAA AGC CTA AAG 231
 His Glu Asp Ile Ile Ser Leu Trp Asp Glu Ser Leu Lys
 65 70 75
 45 CCA TGT GTA AAG TTA ACC CCA CTT TGT GTT ACT CTA AAC 270
 Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn
 80 85 90
 50 TGC AGT GAT GTG AAC AAT TCC ACA AAT CCT AAT GAT ACT 309
 Cys Ser Asp Val Asn Asn Ser Thr Asn Pro Asn Asp Thr
 95 100
 55 AAT ACT AAT TCC ACT AAT ACT ACT TCC TCT ACT CCT ACG 348
 Asn Thr Asn Ser Thr Asn Thr Thr Ser Ser Thr Pro Thr
 105 110 115
 60 GCC ACT ACT AGT AGC GAG GAA AAG ATG GAG AAG GGA GAA 387
 Ala Thr Thr Ser Ser Glu Glu Lys Met Glu Lys Gly Glu
 120 125
 65 ATA AAA AAC TGC TCT TTC AAT ATC ACC ACA CAC ATG AAA 426
 Ile Lys Asn Cys Ser Phe Asn Ile Thr Thr His Met Lys
 130 135 140
 GAT AAG GTA CAG AAA GAA TAT GCA CTT TTT TAT AAA CTT 465
 Asp Lys Val Gln Lys Glu Tyr Ala Leu Phe Tyr Lys Leu
 145 150 155

| | | |
|----|---|------|
| | GAT ATA GTA CCA ATA GAT GAT AAT AAT ACC AGC TAT AGG | 504 |
| | Asp Ile Val Pro Il Asp Asp Asn Asn Thr Ser Tyr Arg | |
| | 160 165 | |
| 5 | TTG ATA AGT TGT AAT ACC TCA GTC ATT ACA CAG GCC TGT | 543 |
| | Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys | |
| | 170 175 180 | |
| 10 | CCA ATG GTG ACC TTT GAG CCA ATT CCC ATA CAT TAT TGT | 582 |
| | Pro Met Val Thr Phe Glu Pro Ile Pro Ile His Tyr Cys | |
| | 185 190 | |
| 15 | GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT AAA GAT AAG | 621 |
| | Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Lys Asp Lys | |
| | 195 200 205 | |
| 20 | AAG TTC AAT GGA ACA GGA CCA TGT TCA AAG GTC AGC ACA | 660 |
| | Lys Phe Asn Gly Thr Gly Pro Cys Ser Lys Val Ser Thr | |
| | 210 215 220 | |
| | GTA CAA TGT ACA CAT GGA ATT AGG CCA GTA GTA TCA ACT | 699 |
| | Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr | |
| | 225 230 | |
| 25 | CAA CTG TTG TTA AAT GGC ACT CTT GCA GAA GAA GAA GTA | 738 |
| | Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val | |
| | 235 240 245 | |
| 30 | GTA ATT AGA TCT GTC AAT TTC ACA CAC AAT GCT AAA ATC | 777 |
| | Val Ile Arg Ser Val Asn Phe Thr Asp Asn Ala Lys Ile | |
| | 250 255 | |
| 35 | ATA ATA GTA CAG CTG AAA GAA CCT GTA GCA ATT AAT TGT | 816 |
| | Ile Ile Val Gln Leu Lys Glu Pro Val Ala Ile Asn Cys | |
| | 260 265 270 | |
| 40 | ACA AGA CCC AAC AAC AAT ACA AGA AAA GGT ATA CAT CTA | 855 |
| | Thr Arg Pro Asn Asn Asn Thr Arg Lys Gly Ile His Leu | |
| | 275 280 285 | |
| | GGA CCA GGG AGC ACA TTT TAT ACA ACA GGA GAA ATA ATA | 894 |
| | Gly Pro Gly Ser Thr Phe Tyr Thr Thr Gly Glu Ile Ile | |
| | 290 295 | |
| 45 | GGA GAC ATA AGA AAA GCA TAT TGC AAC ATT AGT AAA GAA | 933 |
| | Gly Asp Ile Arg Lys Ala Tyr Cys Lys Ile Ser Lys Glu | |
| | 300 305 310 | |
| 50 | AAA TGG AAT AAC ACT TTA AGA CAG GTA GTT AAA AAA TTA | 972 |
| | Lys Trp Asn Asn Thr Leu Arg Gln Val Val Lys Lys Leu | |
| | 315 320 | |
| 55 | AGA GAA CAA TTT GGG AAT AAA ACA ATA ATT TTT AAT CGA | 1011 |
| | Arg Glu Gln Phe Gly Asn Lys Thr Ile Ile Phe Asn Arg | |
| | 325 330 335 | |
| 60 | TCC TCA GGA GGG GAC CCA GAA ATT CTA ATG CAC AGT TTT | 1050 |
| | Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Ser Phe | |
| | 340 345 350 | |
| | AAC TGT GGA GGG GAG TTT TTC TAC TGT AAT ACA ACA CAA | 1089 |
| | Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr Gln | |
| | 355 360 | |

CTG TTT AAT AGT ACT TGG AAT AAT ACT GAA GGG ACA AAT 1128
 Leu Phe Asn Ser Thr Trp Asn Asn Thr Glu Gly Thr Asn
 365 370 375

5 AGC ACT GAA GGA AAT AGC ACA ATC ACA CTC CCA TGC AGA 1167
 Ser Thr Glu Gly Asn Ser Thr Ile Thr Leu Pro Cys Arg
 380 385

10 ATA AAA CAA ATT ATA AAT ATG TGG CAG GAA GTA GGA AAA 1206
 Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys
 390 395 400

15 GCA ACG TAT GCC CCT CCC ATC AGA GGA CGA ATT AGA TGC 1245
 Ala Thr Tyr Ala Pro Pro Ile Arg Gly Arg Ile Arg Cys
 405 410 415

20 ATA TCA AAT ATT ACA GGA CTG CTA TTA ACA AGA GAT GGT 1284
 Ile Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly
 420 425

GGT AGG AAT GTC ACA AAC AAT ACC GAN NCC TTC AGA CCT 1323
 Gly Arg Asn Val Thr Asn Asn Thr Xaa Xaa Phe Arg Pro
 430 435 440

25 GCA GGA GGA GAC ATG AGG GAC AAT TGG AGA AGT GAA TTA 1362
 Gly Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu
 445 450

30 TAT AAA TAT AAA GTA GTA AAA GTT GAA CCA TTA GGA ATA 1401
 Tyr Lys Tyr Lys Val Val Lys Val Glu Pro Leu Gly Ile
 455 460 465

35 GCA CCC ACC AAG GCA AAG AGA AGA GTG GTC CAC AGA GAC 1440
 Ala Pro Thr Lys Ala Lys Arg Arg Val Val His Arg Asp
 470 475 480

40 AAA AGA GCA GCA CTA GGA GCT TTG TTC CTT GGG TTC TTA 1479
 Lys Arg Ala Ala Leu Gly Ala Leu Phe Leu Gly Phe Leu
 485 490

GGA GCA TAA AAG CTT CTA GA 1499
 Gly Ala Xaa Lys Leu Leu
 495 499

45 (2) INFORMATION FOR SEQ ID NO:20:
 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 499 amino acids
 (B) TYPE: Amino Acid
 (D) TOPOLOGY: Linear

50 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:20:
 Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe Cys Ala
 1 5 10 15
 55 Ser Asp Ala Lys Ala Tyr Asp Thr Glu Val His Asn Val Trp Ala
 20 25 30
 Thr His Ala Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Ile Glu
 35 40 45
 60 Leu Val Asn Val Thr Glu Asp Phe Asn Met Trp Lys Asn Lys Met
 50 55 60
 65 Val Asp Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp Glu Ser
 65 70 75

| | | | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|--|
| | Leu | Lys | Pro | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Val | Thr | Leu | Asn | |
| | | | | | 80 | | | | | 85 | | | | | 90 | |
| 5 | Cys | Ser | Asp | Val | Asn | Asn | Ser | Thr | Asn | Pro | Asn | Asp | Thr | Asn | Thr | |
| | | | | | 95 | | | | | 100 | | | | | 105 | |
| | Asn | Ser | Thr | Asn | Thr | Thr | Ser | Ser | Thr | Pro | Thr | Ala | Thr | Thr | Ser | |
| | | | | | 110 | | | | | 115 | | | | | 120 | |
| 10 | Ser | Glu | Glu | Lys | Met | Glu | Lys | Gly | Glu | Ile | Lys | Asn | Cys | Ser | Phe | |
| | | | | | 125 | | | | | 130 | | | | | 135 | |
| | Asn | Ile | Thr | Thr | His | Met | Lys | Asp | Lys | Val | Gln | Lys | Glu | Tyr | Ala | |
| | | | | | 140 | | | | | 145 | | | | | 150 | |
| 15 | Leu | Phe | Tyr | Lys | Leu | Asp | Ile | Val | Pro | Ile | Asp | Asp | Asn | Asn | Thr | |
| | | | | | 155 | | | | | 160 | | | | | 165 | |
| 20 | Ser | Tyr | Arg | Leu | Ile | Ser | Cys | Asn | Thr | Ser | Val | Ile | Thr | Gln | Ala | |
| | | | | | 170 | | | | | 175 | | | | | 180 | |
| | Cys | Pro | Met | Val | Thr | Phe | Glu | Pro | Ile | Pro | Ile | His | Tyr | Cys | Ala | |
| | | | | | 185 | | | | | 190 | | | | | 195 | |
| 25 | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Lys | Asp | Lys | Lys | Phe | Asn | |
| | | | | | 200 | | | | | 205 | | | | | 210 | |
| | Gly | Thr | Gly | Pro | Cys | Ser | Lys | Val | Ser | Thr | Val | Gln | Cys | Thr | His | |
| | | | | | 215 | | | | | 220 | | | | | 225 | |
| 30 | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | Gln | Leu | Leu | Leu | Asn | Gly | Ser | |
| | | | | | 230 | | | | | 235 | | | | | 240 | |
| 35 | Leu | Ala | Glu | Glu | Glu | Val | Val | Ile | Arg | Ser | Val | Asn | Phe | Thr | Asp | |
| | | | | | 245 | | | | | 250 | | | | | 255 | |
| | Asn | Ala | Lys | Ile | Ile | Ile | Val | Gln | Leu | Lys | Glu | Pro | Val | Ala | Ile | |
| | | | | | 260 | | | | | 265 | | | | | 270 | |
| 40 | Asn | Cys | Thr | Arg | Pro | Asn | Asn | Asn | Thr | Arg | Lys | Gly | Ile | His | Leu | |
| | | | | | 275 | | | | | 280 | | | | | 285 | |
| | Gly | Pro | Gly | Ser | Thr | Phe | Tyr | Thr | Thr | Gly | Glu | Ile | Ile | Gly | Asp | |
| | | | | | 290 | | | | | 295 | | | | | 300 | |
| 45 | Ile | Arg | Lys | Ala | Tyr | Cys | Lys | Ile | Ser | Lys | Glu | Lys | Trp | Asn | Asn | |
| | | | | | 305 | | | | | 310 | | | | | 315 | |
| 50 | Thr | Leu | Arg | Gln | Val | Val | Lys | Lys | Leu | Arg | Glu | Gln | Phe | Gly | Asn | |
| | | | | | 320 | | | | | 325 | | | | | 330 | |
| | Lys | Thr | Ile | Ile | Phe | Asn | Arg | Ser | Ser | Gly | Gly | Asp | Pro | Glu | Ile | |
| | | | | | 335 | | | | | 340 | | | | | 345 | |
| 55 | Val | Met | His | Ser | Phe | Asn | Cys | Gly | Gly | Glu | Phe | Phe | Tyr | Cys | Asn | |
| | | | | | 350 | | | | | 355 | | | | | 360 | |
| | Thr | Thr | Gln | Leu | Phe | Asn | Ser | Thr | Trp | Asn | Asn | Thr | Glu | Gly | Thr | |
| | | | | | 365 | | | | | 370 | | | | | 375 | |
| 60 | Asn | Ser | Thr | Glu | Gly | Asn | Ser | Thr | Ile | Thr | Leu | Pro | Cys | Arg | Ile | |
| | | | | | 380 | | | | | 385 | | | | | 390 | |
| 65 | Lys | Gln | Ile | Ile | Asn | Met | Trp | Gln | Glu | Val | Gly | Lys | Ala | Thr | Tyr | |
| | | | | | 395 | | | | | 400 | | | | | 405 | |

Ala Pro Pro Ile Arg Gly Arg Il Arg Cys Ile Ser Asn Ile Thr
410 415 420

5 Gly Leu Leu Leu Thr Arg Asp Gly Gly Arg Asn Val Thr Asn Asn
425 430 435

Thr Xaa Xaa Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp
440 445 450

10 Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Val Glu Pro Leu
455 460 465

Gly Ile Ala Pro Thr Lys Ala Lys Arg Arg Val Val His Arg Asp
470 475 480

15 Lys Arg Ala Ala Leu Gly Ala Leu Phe Leu Gly Phe Leu Gly Ala
485 490 495

20 Xaa Lys Leu Leu
499

(2) INFORMATION FOR SEQ ID NO:21:

(i) SEQUENCE CHARACTERISTICS:

- 25 (A) LENGTH: 1475 base pairs
(B) TYPE: Nucleic Acid
(C) STRANDEDNESS: Single
(D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:21:

30 G GTA CCT GTG TGG AAA GAA GCA AAC ACA ACT CTA TTT 37
Val Pro Val Trp Lys Glu Ala Asn Thr Thr Leu Phe
1 5 10

35 TGT GCA TCA GAT GCT AAA GCA TAT GAT AGA GAA GTA CAT 76
Cys Ala Ser Asp Ala Lys Ala Tyr Asp Arg Glu Val His
15 20 25

40 AAT GTT TGG GCA ACA CAT GCC TGT GTA CCC ACA GAC CCC 115
Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro
30 35

AAC CCA CAA GAA ATA GTA TTG GGA AAT GTG ACA GAA AAT 154
Asn Pro Gln Glu Ile Val Leu Gly Asn Val Thr Glu Asn
40 45 50

45 TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA ATG CAT 193
Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His
55 60

50 GAG GAT ATA ATC AAT TTA TGG GAT CAA AGC TTA AAG CCA 232
Glu Asp Ile Ile Asn Leu Trp Asp Gln Ser Leu Lys Pro
65 70 75

55 TGT GTA AAG TTA ACT CCA CTC TGT GTT ACT TTA AAG TGC 271
Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Lys Cys
80 85 90

60 AAG GAT CTG GAG AGG AAT ACT ACC TAT AAT AGC ACT ATT 310
Lys Asp Leu Glu Arg Asn Thr Thr Tyr Asn Ser Thr Ile
95 100

65 ACC AAT AAT AGT AGT TTG GAG GGA CTA AGA GAA CAA ATG 349
Thr Asn Asn Ser Ser Leu Glu Gly Leu Arg Glu Gln Met
105 110 115

| | |
|----|---|
| | ACA AAC TGC TCT TTC AAC ATC ACC ACA AGT ATA AGA GAT 388 |
| | Thr Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Arg Asp |
| | 120 125 |
| 5 | AAG GTG CAG AAA GAA TAT GCA CTT TTG TAT AAA CTT GAT 427 |
| | Lys Val Gln Lys Glu Tyr Ala Leu Leu Tyr Lys Leu Asp |
| | 130 135 140 |
| 10 | GTA GTA CCA ATA GAA GAA GAT GAC AAT ACT AGC TAT AGA 466 |
| | Val Val Pro Ile Glu Glu Asp Asp Asn Thr Ser Tyr Arg |
| | 145 150 155 |
| 15 | TTG ATA AGT TGT AAC ACC TCA GTC ATT ACA CAG GCT TGT 505 |
| | Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys |
| | 160 165 |
| 20 | CCA AAG ACA TCC TTT GAG CCA ATT CCC ATA CAT TAT TGT 544 |
| | Pro Lys Thr Ser Phe Glu Pro Ile Pro Ile His Tyr Cys |
| | 170 175 180 |
| 25 | GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT AAT GAT AAG 583 |
| | Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asp Lys |
| | 185 190 |
| 30 | AAG TTC AAT GGA ACA GGA CCA TGT AAA AAT CTC ACC ACA 622 |
| | Lys Phe Asn Gly Thr Gly Pro Cys Lys Asn Val Ser Thr |
| | 195 200 205 |
| 35 | GTA CAA TGT ACA CAT GGA ATT AGG CCA GTA GTA TCA ACT 661 |
| | Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr |
| | 210 215 220 |
| 40 | CAA CTG TTG TTA AAT GGC AGT CTA GCA GAA GAA GAG GTA 700 |
| | Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val |
| | 225 230 |
| 45 | GTA ATC AGA TCT GCC AAT TTC ACA GAC AAT GCT AAA ACC 739 |
| | Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr |
| | 235 240 245 |
| 50 | ATA ATA GTA CAT CTA AAT GAA ACT GTA AAA ATT AAT TGT 778 |
| | Ile Ile Val His Leu Asn Glu Thr Val Lys Ile Asn Cys |
| | 250 255 |
| 55 | ACA AGA CTT GGC AAC AAT ACA AGA AAA AGT ATA AAT ATA 817 |
| | Thr Arg Leu Gly Asn Asn Thr Arg Lys Ser Ile Asn Ile |
| | 260 265 270 |
| 60 | GGA CCA GGG AGA GTA CTC TAT GCA ACA GGA GAA ATA ATA 856 |
| | Gly Pro Gly Arg Val Leu Tyr Ala Thr Gly Glu Ile Ile |
| | 275 280 285 |
| 65 | GGA GAC ATA AGA CAA GCA CAT TGT AAC ATT AGT AGA GCA 895 |
| | Gly Asp Ile Arg Gln Ala His Cys Asn Ile Ser Arg Ala |
| | 290 295 |
| 70 | CAA TGG AAT AAG ACT TTA GAA AAG GTA GTT GAC AAA TTA 934 |
| | Gln Trp Asn Lys Thr Leu Glu Lys Val Val Asp Lys Leu |
| | 300 305 310 |
| 75 | AGA AAA CAA TTT GGG GAT AAT ACA ACA ATA GCT TTT AAT 973 |
| | Arg Lys Gln Phe Gly Asp Asn Thr Thr Ile Ala Phe Asn |
| | 315 320 |

CGA TCC TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC ACT 1012
 Arg Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Thr
 325 330 335

5 TTT AAT TGT GGA GGG GAA TTT TTC TAC TGT AAT ACA ACA 1051
 Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr
 340 345 350

10 CAA CTG TTT AAT AGT ACT TGG AAT AAT ACT TGG AAG GAT 1090
 Gln Leu Phe Asn Ser Thr Trp Asn Asn Thr Trp Lys Asp
 355 360

15 CCT AAC AGG AGT GAC AAT ATC ACA CTC CCA TGC AGA ATA 1129
 Pro Asn Arg Ser Asp Asn Ile Thr Leu Pro Cys Arg Ile
 365 370 375

20 AAA CAA ATT ATA AAC ATG TGG CAG GAA GTA GGA AAA GCA 1168
 Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala
 380 385

25 ATG TAC GCC CCT CCC ATC AGA GGG GAA ATT AGA TGT TCA 1207
 Met Tyr Ala Pro Pro Ile Arg Gly Glu Ile Arg Cys Ser
 390 395 400

30 AAT GAC GAT GGT AAT GAC ACG ACC ACA AAC AGG ACC GAG 1285
 Asn Asp Asp Gly Asn Asp Thr Thr Thr Asn Arg Thr Glu
 420 425

35 ATC TTC AGA CCT GGA GGA GGA GAT ATG AGG GAC AAT TGG 1324
 Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp
 430 435 440

40 AGA AGT GAA TTA TAT AGA TAT AAA GTA GTA AAA ATT GAA 1363
 Arg Ser Glu Leu Tyr Arg Tyr Lys Val Val Lys Ile Glu
 445 450

45 CCA TTA GGA ATA GCA CCC ACC AGG GCA AAG AGA AGA GTG 1402
 Pro Leu Gly Ile Ala Pro Thr Arg Ala Lys Arg Arg Val
 455 460 465

50 GTG CAG AGA GAA AAA AGA GCA GTA GGA CTA GGA GCT TTG 1441
 Val Gln Arg Glu Lys Arg Ala Val Gly Leu Gly Ala Leu
 470 475 480

TTC CTT GGG T TCTTAGGAG CATAAAGCTT CTAGA 1475
 Phe Leu Gly
 483

(2) INFORMATION FOR SEQ ID NO:22:
 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 491 amino acids
 (B) TYPE: Amino Acid
 (D) TOPOLOGY: Linear
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:22:

60 Val Pro Val Trp Lys Glu Ala Asn Thr Thr Leu Phe Cys Ala Ser
 1 5 10 15

65 Asp Ala Lys Ala Tyr Asp Arg Glu Val His Asn Val Trp Ala Thr
 20 25 30

| | | | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|
| | His | Ala | Cys | Val | Pro | Thr | Asp | Pro | Asn | Pro | Gln | Glu | Ile | Val | Leu | |
| | | | | | 35 | | | | | 40 | | | | | | 45 |
| 5 | Gly | Asn | Val | Thr | Glu | Asn | Phe | Asn | Met | Trp | Lys | Asn | Asn | Met | Val | |
| | | | | | 50 | | | | | 55 | | | | | | 60 |
| | Glu | Gln | Met | His | Glu | Asp | Ile | Ile | Asn | Leu | Trp | Asp | Gln | Ser | Leu | |
| | | | | | 65 | | | | | 70 | | | | | | 75 |
| 10 | Lys | Pro | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Val | Thr | Leu | Lys | Cys | |
| | | | | | 80 | | | | | 85 | | | | | | 90 |
| | Lys | Asp | Leu | Glu | Arg | Asn | Thr | Thr | Tyr | Asn | Ser | Thr | Ile | Thr | Asn | |
| | | | | | 95 | | | | | 100 | | | | | | 105 |
| 15 | Asn | Ser | Ser | Leu | Glu | Gly | Leu | Arg | Glu | Gln | Met | Thr | Asn | Cys | Ser | |
| | | | | | 110 | | | | | 115 | | | | | | 120 |
| | Phe | Asn | Ile | Thr | Thr | Ser | Ile | Arg | Asp | Lys | Val | Gln | Lys | Glu | Tyr | |
| | | | | | 125 | | | | | 130 | | | | | | 135 |
| 20 | Ala | Leu | Leu | Tyr | Lys | Leu | Asp | Val | Val | Pro | Ile | Glu | Glu | Asp | Asp | |
| | | | | | 140 | | | | | 145 | | | | | | 150 |
| 25 | Asn | Thr | Ser | Tyr | Arg | Leu | Ile | Ser | Cys | Asn | Thr | Ser | Val | Ile | Thr | |
| | | | | | 155 | | | | | 160 | | | | | | 165 |
| | Gln | Ala | Cys | Pro | Lys | Thr | Ser | Phe | Glu | Pro | Ile | Pro | Ile | His | Tyr | |
| | | | | | 170 | | | | | 175 | | | | | | 180 |
| 30 | Cys | Ala | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Asn | Asp | Lys | Lys | |
| | | | | | 185 | | | | | 190 | | | | | | 195 |
| | Phe | Asn | Gly | Thr | Gly | Pro | Cys | Lys | Asn | Val | Ser | Thr | Val | Gln | Cys | |
| | | | | | 200 | | | | | 205 | | | | | | 210 |
| 35 | Thr | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | Gln | Leu | Leu | Leu | Asn | |
| | | | | | 215 | | | | | 220 | | | | | | 225 |
| 40 | Gly | Ser | Leu | Ala | Glu | Glu | Glu | Val | Val | Ile | Arg | Ser | Ala | Asn | Phe | |
| | | | | | 230 | | | | | 235 | | | | | | 240 |
| | Thr | Asp | Asn | Ala | Lys | Thr | Ile | Ile | Val | His | Leu | Asn | Glu | Thr | Val | |
| | | | | | 245 | | | | | 250 | | | | | | 255 |
| 45 | Lys | Ile | Asn | Cys | Thr | Arg | Leu | Gly | Asn | Asn | Thr | Arg | Lys | Ser | Ile | |
| | | | | | 260 | | | | | 265 | | | | | | 270 |
| | Asn | Ile | Gly | Pro | Gly | Arg | Val | Leu | Tyr | Ala | Thr | Gly | Glu | Ile | Ile | |
| | | | | | 275 | | | | | 280 | | | | | | 285 |
| 50 | Gly | Asp | Ile | Arg | Gln | Ala | His | Cys | Asn | Ile | Ser | Arg | Ala | Gln | Trp | |
| | | | | | 290 | | | | | 295 | | | | | | 300 |
| 55 | Asn | Lys | Thr | Leu | Glu | Lys | Val | Val | Asp | Lys | Leu | Arg | Lys | Gln | Phe | |
| | | | | | 305 | | | | | 310 | | | | | | 315 |
| | Gly | Asp | Asn | Thr | Thr | Ile | Ala | Phe | Asn | Arg | Ser | Ser | Gly | Gly | Asp | |
| | | | | | 320 | | | | | 325 | | | | | | 330 |
| 60 | Pro | Glu | Ile | Val | Met | His | Thr | Phe | Asn | Cys | Gly | Gly | Glu | Phe | Phe | |
| | | | | | 335 | | | | | 340 | | | | | | 345 |
| | Tyr | Cys | Asn | Thr | Thr | Gln | Leu | Phe | Asn | Ser | Thr | Trp | Asn | Asn | Thr | |
| | | | | | 350 | | | | | 355 | | | | | | 360 |

Trp Lys Asp Pro Asn Arg Ser Asp Asn Ile Thr Leu Pro Cys Arg
 365 370 375
 5 Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met
 380 385 390
 Tyr Ala Pro Pro Ile Arg Gly Glu Ile Arg Cys Ser Ser Asn Ile
 395 400 405
 10 Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Asp Asp Gly Asn
 410 415 420
 Asp Thr Thr Thr Asn Arg Thr Glu Ile Phe Arg Pro Gly Gly Gly
 425 430 435
 15 Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr Arg Tyr Lys Val
 440 445 450
 Val Lys Ile Glu Pro Leu Gly Ile Ala Pro Thr Arg Ala Lys Arg
 455 460 465
 20 Arg Val Val Gln Arg Glu Lys Arg Ala Val Gly Leu Gly Ala Leu
 470 475 480
 25 Phe Leu Gly Phe Leu Gly Ala Leu Phe Leu Gly
 485 490 491

(2) INFORMATION FOR SEQ ID NO:23:

(i) SEQUENCE CHARACTERISTICS:

- 30 (A) LENGTH: 1475 base pairs
 (B) TYPE: Nucleic Acid
 (C) STRANDEDNESS: Single
 (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:23:

35 G GTA CCT GTG TGG AAA GAA GCA AAC ACA ACT CTA TTT 37
 Val Pro Val Trp Lys Glu Ala Asn Thr Thr Leu Phe
 1 5 10
 40 TGT GCA TCA GAT GCT AAA GCA TAT GAT AGA GAA GTA CAT 76
 Cys Ala Ser Asp Ala Lys Ala Tyr Asp Arg Glu Val His
 15 20 25
 45 AAT GTT TGG GCA ACA CAT GCC TGT GTA CCC ACA GAC CCC 115
 Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro
 30 35
 50 AAC CCA CAA GAA ATA GTA TTG GGA AAT GTG ACA GAA AAT 154
 Asn Pro Gln Glu Ile Val Leu Gly Asn Val Thr Glu Asn
 40 45 50
 TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAA ATG CAT 193
 Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His
 55 60
 55 GAG GAT ATA ATC AAT TTA TGG GAT CAA AGC TTA AAG CCA 232
 Glu Asp Ile Ile Asn Leu Trp Asp Gln Ser Leu Lys Pro
 65 70 75
 60 TGT GTA AAG TTA ACT CCA CTC TGT GTT ACT TTA AAG TGC 271
 Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Lys Cys
 80 85 90

AAG GAT CTG GAG AGG AAT ACT ACC TAT AAT AGC ACT ATT 310
 Lys Asp Leu Glu Arg Asn Thr Thr Tyr Asn Ser Thr Ile
 95 100

5 ACC AAT AAT AGT AGT TTG GAG GGA CTA AGA GAA CAA ATG 349
 Thr Asn Asn Ser Ser Leu Glu Gly Leu Arg Glu Gln Met
 105 110 115

10 ACA AAC TGC TCT TTC AAC ATC ACC ACA AGT ATA AGA GAT 388
 Thr Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Arg Asp
 120 125

15 AAG GTG CAG AAA GAA TAT GCA CTT TTG TAT AAA CTT GAT 427
 Lys Val Gln Lys Glu Tyr Ala Leu Leu Tyr Lys Leu Asp
 130 135 140

20 GTA GTA CCA ATA GAA GAA GAT GAC AAT ACT AGC TAT AGA 466
 Val Val Pro Ile Glu Glu Asp Asp Asn Thr Ser Tyr Arg
 145 150 155

TTG ATA AGT TGT AAC ACC TCA GTC ATT ACA CAG GCT TGT 505
 Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys
 160 165

25 CCA AAG ACA TCC TTT GAG CCA ATT CCC ATA CAT TAT TGT 544
 Pro Lys Thr Ser Phe Glu Pro Ile Pro Ile His Tyr Cys
 170 175 180

30 GCC CCG GCT GGT TTT GCG ATT CTA AAG TGT AAT GAT AAG 583
 Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asp Lys
 185 190

35 AAG TTC AAT GGA ACA GGA CCA TGT AAA AAT GTC ACC ACA 622
 Lys Phe Asn Gly Thr Gly Pro Cys Lys Asn Val Ser Thr
 195 200 205

40 GTA CAA TGT ACA CAT GGA ATT AGG CCA GTA GTA TCA ACT 661
 Val Gln Cys Thr His Gly Ile Arg Pro Val Val Ser Thr
 210 215 220

CAA CTG TTG TTA AAT GGC AGT CTA GCA GAA GAA GAG GTA 700
 Gln Leu Leu Leu Asn Gly Ser Leu Ala Glu Glu Glu Val
 225 230

45 GTA ATC AGA TCT GCC AAT TTC ACA GAC AAT GCT AAA ACC 739
 Val Ile Arg Ser Ala Asn Phe Thr Asp Asn Ala Lys Thr
 235 240 245

50 ATA ATA GTA CAT CTA AAT GAA ACT GTA AAA ATT AAT TGT 778
 Ile Ile Val His Leu Asn Glu Thr Val Lys Ile Asn Cys
 250 255

55 ACA AGA CTT GGC AAC AAT ACA AGA AAA AGT ATA AAT ATA 817
 Thr Arg Leu Gly Asn Asn Thr Arg Lys Ser Ile Asn Ile
 260 265 270

60 GGA CCA GGG AGA GTA CTC TAT GCA ACA GGA GAA ATA ATA 856
 Gly Pro Gly Arg Val Leu Tyr Ala Thr Gly Glu Ile Ile
 275 280 285

GGA GAC ATA AGA CAA GCA CAT TGT AAC ATT AGT AGA GCA 895
 Gly Asp Ile Arg Gln Ala His Cys Asn Ile Ser Arg Ala
 290 295

CAA TGG AAT AAG ACT TTA GAA AAG GTA GTT GAC AAG TTA 934
 Gln Trp Asn Lys Thr Leu Glu Lys Val Val Asp Lys Leu
 300 305 310

5 AGA AAA CAA TTT GGG GAT AAT ACA ACA ATA GCT TTT AAT 973
 Arg Lys Gln Phe Gly Asp Asn Thr Thr Ile Ala Phe Asn
 315 320

10 CGA TCC TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC ACT 1012
 Arg Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Thr
 325 330 335

15 TTT AAT TGT GGA GGG GAA TTT TTC TAC TGT AAT ACA ACA 1051
 Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr
 340 345 350

20 CAA CTG TTT AAT AGT ACT TGG AAT AAT ACT TGG AAG GAT 1090
 Gln Leu Phe Asn Thr Trp Asn Asn Thr Trp Lys Asp
 355 360

CCT AAC AGG AGT GAC AAT ATC ACA CTC CCA TGC AGA ATA 1129
 Pro Asn Arg Ser Asp Asn Ile Thr Leu Pro Cys Arg Ile
 365 370 375

25 AAA CAA ATT ATA AAC ATG TGG CAG GAA GTA GGA AAA GCA 1168
 Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala
 380 385

30 ATG TAC GCC CCT CCC ATC AGA GGG GAA ATT AGA TGT TCA 1207
 Met Tyr Ala Pro Pro Ile Arg Gly Glu Ile Arg Cys Ser
 390 395 400

35 TCA AAT ATC ACA GGG CTG CTA CTA ACA AGA GAT GGT GGT 1246
 Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 405 410 415

40 AAT GAC GAT GGT AAT GAC ACG ACC ACA AAC AGG ACC GAG 1285
 Asn Asp Asp Gly Asn Asp Thr Thr Thr Asn Arg Thr Glu
 420 425

ATC TTC AGA CCT GGA GGA GGA GAT ATG AGG GAC AAT TGG 1324
 Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp
 430 435 440

45 AGA AGT GAA TTA TAT AGA TAT AAA GTA GTA AAA ATT GAA 1363
 Arg Ser Glu Leu Tyr Arg Tyr Lys Val Val Lys Ile Glu
 445 450

50 CCA TTA GGA ATA GCA CCC ACC AGG GCA AAG AGA AGA GTG 1402
 Pro Leu Gly Ile Ala Pro Thr Arg Ala Lys Arg Arg Val
 455 460 465

55 GTG CAG AGA GAA AAA AGA GCA GTA GGA CTA GGA GCT TTG 1441
 Val Gln Arg Glu Lys Arg Ala Val Gly Leu Gly Ala Leu
 470 475 480

TTC CTT GGG TTC TTG GGA GCA TAA AGC TTC TAG A 1475
 Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa
 485 490 491

(2) INFORMATION FOR SEQ ID NO:24:

(1) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 491 amino acids

(B) TYPE: Amino Acid

(D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:24:

| | | | | | | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|----|
| | Val | Pro | Val | Trp | Lys | Glu | Ala | Asn | Thr | Thr | Leu | Phe | Cys | Ala | Ser | 1 | 5 | 10 | 15 |
| 5 | Asp | Ala | Lys | Ala | Tyr | Asp | Arg | Glu | Val | His | Asn | Val | Trp | Ala | Thr | 20 | 25 | 30 | |
| 10 | His | Ala | Cys | Val | Pro | Thr | Asp | Pro | Asn | Pro | Gln | Glu | Ile | Val | Leu | 35 | 40 | 45 | |
| | Gly | Asn | Val | Thr | Glu | Asn | Phe | Asn | Met | Trp | Lys | Asn | Asn | Met | Val | 50 | 55 | 60 | |
| 15 | Glu | Gln | Met | His | Glu | Asp | Ile | Ile | Asn | Leu | Trp | Asp | Gln | Ser | Leu | 65 | 70 | 75 | |
| | Lys | Pro | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Val | Thr | Leu | Lys | Cys | 80 | 85 | 90 | |
| 20 | Lys | Asp | Leu | Glu | Arg | Asn | Thr | Thr | Tyr | Asn | Ser | Thr | Ile | Thr | Asn | 95 | 100 | 105 | |
| | Asn | Ser | Ser | Leu | Glu | Gly | Leu | Arg | Glu | Gln | Met | Thr | Asn | Cys | Ser | 110 | 115 | 120 | |
| 25 | Phe | Asn | Ile | Thr | Thr | Ser | Ile | Arg | Asp | Lys | Val | Gln | Lys | Glu | Tyr | 125 | 130 | 135 | |
| 30 | Ala | Leu | Leu | Tyr | Lys | Leu | Asp | Val | Val | Pro | Ile | Glu | Glu | Asp | Asp | 140 | 145 | 150 | |
| | Asn | Thr | Ser | Tyr | Arg | Leu | Ile | Ser | Cys | Asn | Thr | Ser | Val | Ile | Thr | 155 | 160 | 165 | |
| 35 | Gln | Ala | Cys | Pro | Lys | Thr | Ser | Phe | Glu | Pro | Ile | Pro | Ile | His | Tyr | 170 | 175 | 180 | |
| | Cys | Ala | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Asn | Asp | Lys | Lys | 185 | 190 | 195 | |
| 40 | Phe | Asn | Gly | Thr | Gly | Pro | Cys | Lys | Asn | Val | Ser | Thr | Val | Gln | Cys | 200 | 205 | 210 | |
| 45 | Thr | His | Gly | Ile | Arg | Pro | Val | Val | Ser | Thr | Gln | Leu | Leu | Leu | Asn | 215 | 220 | 225 | |
| | Gly | Ser | Leu | Ala | Glu | Glu | Glu | Val | Val | Ile | Arg | Ser | Ala | Asn | Phe | 230 | 235 | 240 | |
| 50 | Thr | Asp | Asn | Ala | Lys | Thr | Ile | Ile | Val | His | Leu | Asn | Glu | Thr | Val | 245 | 250 | 255 | |
| | Lys | Ile | Asn | Cys | Thr | Arg | Leu | Gly | Asn | Asn | Thr | Arg | Lys | Ser | Ile | 260 | 265 | 270 | |
| 55 | Asn | Ile | Gly | Pro | Gly | Arg | Val | Leu | Tyr | Ala | Thr | Gly | Glu | Ile | Ile | 275 | 280 | 285 | |
| 60 | Gly | Asp | Ile | Arg | Gln | Ala | His | Cys | Asn | Ile | Ser | Arg | Ala | Gln | Trp | 290 | 295 | 300 | |
| | Asn | Lys | Thr | Leu | Glu | Lys | Val | Val | Asp | Lys | Leu | Arg | Lys | Gln | Phe | 305 | 310 | 315 | |
| 65 | | | | | | | | | | | | | | | | | | | |

Gly Asp Asn Thr Thr Ile Ala Phe Asn Arg Ser Ser Gly Gly Asp
 320 325 330
 5 Pro Glu Ile Val Met His Thr Phe Asn Cys Gly Gly Glu Phe Phe
 335 340 345
 Tyr Cys Asn Thr Thr Gln Leu Phe Asn Ser Thr Trp Asn Asn Thr
 350 355 360
 10 Trp Lys Asp Pro Asn Arg Ser Asp Asn Ile Thr Leu Pro Cys Arg
 365 370 375
 Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met
 380 385 390
 15 Tyr Ala Pro Pro Ile Arg Gly Glu Ile Arg Cys Ser Ser Asn Ile
 395 400 405
 Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Asp Asp Gly Asn
 410 415 420
 20 Asp Thr Thr Thr Asn Arg Thr Glu Ile Phe Arg Pro Gly Gly Gly
 425 430 435
 25 Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr Arg Tyr Lys Val
 440 445 450
 Val Lys Ile Glu Pro Leu Gly Ile Ala Pro Thr Arg Ala Lys Arg
 455 460 465
 30 Arg Val Val Gln Arg Glu Lys Arg Ala Val Gly Leu Gly Ala Leu
 470 475 480
 Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa
 485 490 491
 35

(2) INFORMATION FOR SEQ ID NO:25:

(i) SEQUENCE CHARACTERISTICS:

- 40 (A) LENGTH: 1435 base pairs
 (B) TYPE: Nucleic Acid
 (C) STRANDEDNESS: Single
 (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:25:

45 CTC GAG GTA CCT GTG TGG AAA GAA GCA ACC ACC ACT 36
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr
 1 5 10
 50 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT GAT TCA GAG 75
 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Ser Glu
 15 20 25
 55 GCA CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA 114
 Ala His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr
 30 35
 GAC CCC AAC CCA CAA GAA GTA GAA TTG GAA AAT GTG ACA 153
 Asp Pro Asn Pro Gln Glu Val Glu Leu Glu Asn Val Thr
 40 45 50
 60 GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAG 192
 Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln
 55 60

| | |
|----|---|
| | ATG CAT GGG GAT ATA ATT AGT TTA TGG GAT CAA AGC CTA 231 |
| | Met His Gly Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu |
| | 65 70 75 |
| 5 | AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT GTT ACG TTA 270 |
| | Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu |
| | 80 85 90 |
| 10 | AAT TGC ACT GAC CCA AAT GTT ACT AAT AGC GAG AGA ACG 309 |
| | Asn Cys Thr Asp Pro Asn Val Thr Asn Ser Glu Arg Thr |
| | 95 100 |
| 15 | ATA GAG GGG GGA GAA ATA AAA AAT TGC TCT TTC AAT ATC 348 |
| | Ile Glu Gly Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile |
| | 105 110 115 |
| 20 | ACC ACA AAC ATA AGA GAT AGG TTT CAG AAA GAA TAT GCA 387 |
| | Thr Thr Asn Ile Arg Asp Arg Phe Gln Lys Glu Tyr Ala |
| | 120 125 |
| | CTT TTT TAT AAA CTT GAT GTA ATA CCA TTA GGT AAT GAT 426 |
| | Leu Phe Tyr Lys Leu Asp Val Ile Pro Leu Gly Asn Asp |
| | 130 135 140 |
| 25 | AAT ACT AGC TAT AGG TTG ATA AGT TGT AAC ACC TCA GTC 465 |
| | Asn Thr Ser Tyr Arg Leu Ile Ser Cys Asn Thr Ser Val |
| | 145 150 155 |
| 30 | ATT ACA CAG GCC TGT CCA AAG GTA TCC TTT GAG CCA ATT 504 |
| | Ile Thr Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile |
| | 160 165 |
| 35 | CCC ATA CAT TAT TGT GCC CCG GCT GGT TTT CCG ATT CTA 543 |
| | Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile Leu |
| | 170 175 180 |
| 40 | AAG TGT AAA GAT AAG AAG TTC AAT GGA ACA GGA CCA TGT 582 |
| | Lys Cys Lys Asp Lys Lys Phe Asn Gly Thr Gly Pro Cys |
| | 185 190 |
| | ACA AAT GTC AGC ACA GTA CAA TGT ACA CAT GGA ATT AAG 621 |
| | Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Lys |
| | 195 200 205 |
| 45 | CCA GTA GTA TCA ACT CAA CTG TTG TTA AAT GGC AGT CTA 660 |
| | Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu |
| | 210 215 220 |
| 50 | GCA GAA GAA GAC ATA GTA ATT AGA TCC GCC AAT CTC ACA 699 |
| | Ala Glu Glu Asp Ile Val Ile Arg Ser Ala Asn Leu Thr |
| | 225 230 |
| 55 | GAC AAT GCT AAA AAC ATA ATA GTA CAG CTG AAT GAA TCT 738 |
| | Asp Asn Ala Lys Asn Ile Ile Val Gln Leu Asn Glu Ser |
| | 235 240 245 |
| 60 | GTA ACA ATG AAT TGT ACA AGA CCC AAC AAC AAT ACA ATG 777 |
| | Val Thr Met Asn Cys Thr Arg Pro Asn Asn Asn Thr Met |
| | 250 255 |
| | AAA AGT ATA CAT ATA GGA CCA GGC ACA GCA TTT TAT GCA 816 |
| | Lys Ser Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala |
| | 260 265 270 |

ACA GGA AAC ATA ATA GGA GAT ATA AGA CAA GCA CAT TGT 855
 Thr Gly Asn Ile Ile Gly Asp Ile Arg Gln Ala His Cys
 275 280 285

5 AAC ATT AGT GGA ACA AAA TGG AAT GAC ACT TTG AAA AAG 894
 Asn Ile Ser Gly Thr Lys Trp Asn Asp Thr Leu Lys Lys
 290 295

10 ATA GCT ATA AAA TTA AGA GAA CAA TTT AAT AAG ACA ATA 933
 Ile Ala Ile Lys Leu Arg Glu Gln Phe Asn Lys Thr Ile
 300 305 310

15 GTC TTT AAT CAA TCC TCA GGA GGG GAC CCA GAA ATT GCA 972
 Val Phe Asn Gln Ser Ser Gly Gly Asp Pro Glu Ile Ala
 315 320

20 ACG CTC AGT TTT AAT TGT GGA GGG GAA TTT TTC TAC TGT 1011
 Thr Leu Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys
 325 330 335

AAT TCA ACA CAA CTG TTT AAT AGT ACT TGG AAT AGT ACT 1050
 Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Asn Ser Thr
 340 345 350

25 GGG TCA AAT AAC ACT AAA GGA AAT GAC ACA ATC ACA CTC 1089
 Gly Ser Asn Asn Thr Lys Gly Asn Asp Thr Ile Thr Leu
 355 360

30 CCA TGC AGA ATA AGA CAA ATT ATA AAC ATG TGG CAG AAA 1128
 Pro Cys Arg Ile Arg Gln Ile Ile Asn Met Trp Gln Lys
 365 370 375

35 ATA GGA AAA GCA ATG TAT GCC CCT CCC ATC AAA GGG CAA 1167
 Ile Gly Lys Ala Met Tyr Ala Pro Pro Ile Lys Gly Gln
 380 385

40 ATT AGA TGT TCA TCA AAT ATT ACA GGG CTA ATA TTA ACA 1206
 Ile Arg Cys Ser Ser Asn Ile Thr Gly Leu Ile Leu Thr
 390 395 400

AGA GAT GGT GGT AAC AAC AAC ATG AGC AAG ACC ACC GAG 1245
 Arg Asp Gly Gly Asn Asn Asn Met Ser Lys Thr Thr Glu
 405 410 415

45 ACC TTC AGA CCT GGA GGA GGA GAT ATG AGG GAC AAT TGG 1284
 Thr Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp
 420 425

50 AGA AGT GAA TTA TAT AAA TAT AAA GTA GTA AAA ATT GAA 1323
 Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu
 430 435 440

55 CCA TTA GGA GTA GCA CCC ACC AGG GCA AAG AGA AGA GTG 1362
 Pro Leu Gly Val Ala Pro Thr Arg Ala Lys Arg Arg Val
 445 450

60 GTG CAG AGA GAA AAA AGA GCA GTG GGA ATA GGA GCT GTG 1401
 Val Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Val
 455 460 465

TTC CTT GGG TTC TTG GGA GCA TAA AGC TTC TAG A 1435
 Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa
 470 475 478

(2) INFORMATION FOR SEQ ID NO:26:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 478 amino acids

(B) TYPE: Amino Acid

(D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:26:

| | | | | | | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|----|
| 5 | Leu | Glu | Val | Pro | Val | Trp | Lys | Glu | Ala | Thr | Thr | Thr | Leu | Phe | Cys | 1 | 5 | 10 | 15 |
| 10 | Ala | Ser | Asp | Ala | Lys | Ala | Tyr | Asp | Ser | Glu | Ala | His | Asn | Val | Trp | 20 | 25 | 30 | |
| 15 | Ala | Thr | His | Ala | Cys | Val | Pro | Thr | Asp | Pro | Asn | Pro | Gln | Glu | Val | 35 | 40 | 45 | |
| | Glu | Leu | Glu | Asn | Val | Thr | Glu | Asn | Phe | Asn | Met | Trp | Lys | Asn | Asn | 50 | 55 | 60 | |
| 20 | Met | Val | Glu | Gln | Met | His | Gly | Asp | Ile | Ile | Ser | Leu | Trp | Asp | Gln | 65 | 70 | 75 | |
| | Ser | Leu | Lys | Pro | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Val | Thr | Leu | 80 | 85 | 90 | |
| 25 | Asn | Cys | Thr | Asp | Pro | Asn | Val | Thr | Asn | Ser | Glu | Arg | Thr | Ile | Glu | 95 | 100 | 105 | |
| 30 | Gly | Gly | Glu | Ile | Lys | Asn | Cys | Ser | Phe | Asn | Ile | Thr | Thr | Asn | Ile | 110 | 115 | 120 | |
| | Arg | Asp | Arg | Phe | Gln | Lys | Glu | Tyr | Ala | Leu | Phe | Tyr | Lys | Leu | Asp | 125 | 130 | 135 | |
| 35 | Val | Ile | Pro | Leu | Gly | Asn | Asp | Asn | Thr | Ser | Tyr | Arg | Leu | Ile | Ser | 140 | 145 | 150 | |
| | Cys | Asn | Thr | Ser | Val | Ile | Thr | Gln | Ala | Cys | Pro | Lys | Val | Ser | Phe | 155 | 160 | 165 | |
| 40 | Glu | Pro | Ile | Pro | Ile | His | Tyr | Cys | Ala | Pro | Ala | Gly | Phe | Ala | Ile | 170 | 175 | 180 | |
| 45 | Leu | Lys | Cys | Lys | Asp | Lys | Lys | Phe | Asn | Gly | Thr | Gly | Pro | Cys | Thr | 185 | 190 | 195 | |
| | Asn | Val | Ser | Thr | Val | Gln | Cys | Thr | His | Gly | Ile | Lys | Pro | Val | Val | 200 | 205 | 210 | |
| 50 | Ser | Thr | Gln | Leu | Leu | Leu | Asn | Gly | Ser | Leu | Ala | Glu | Glu | Asp | Ile | 215 | 220 | 225 | |
| | Val | Ile | Arg | Ser | Ala | Asn | Leu | Thr | Asp | Asn | Ala | Lys | Asn | Ile | Ile | 230 | 235 | 240 | |
| 55 | Val | Gln | Leu | Asn | Glu | Ser | Val | Thr | Met | Asn | Cys | Thr | Arg | Pro | Asn | 245 | 250 | 255 | |
| 60 | Asn | Asn | Thr | Met | Lys | Ser | Ile | His | Ile | Gly | Pro | Gly | Arg | Ala | Phe | 260 | 265 | 270 | |
| | Tyr | Ala | Thr | Gly | Asn | Ile | Ile | Gly | Asp | Ile | Arg | Gln | Ala | His | Cys | 275 | 280 | 285 | |

Asn Ile Ser Gly Thr Lys Trp Asn Asp Thr Leu Lys Lys Ile Ala
 290 295 300
 5 Ile Lys Leu Arg Glu Gln Phe Asn Lys Thr Ile Val Phe Asn Gln
 305 310 315
 Ser Ser Gly Gly Asp Pro Glu Ile Ala Thr Leu Ser Phe Asn Cys
 320 325 330
 10 Gly Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser
 335 340 345
 Thr Trp Asn Ser Thr Gly Ser Asn Asn Thr Lys Gly Asn Asp Thr
 350 355 360
 15 Ile Thr Leu Pro Cys Arg Ile Arg Gln Ile Ile Asn Met Trp Gln
 365 370 375
 Lys Ile Gly Lys Ala Met Tyr Ala Pro Pro Ile Lys Gly Gln Ile
 380 385 390
 20 Arg Cys Ser Ser Asn Ile Thr Gly Leu Ile Leu Thr Arg Asp Gly
 395 400 405
 25 Gly Asn Asn Asn Met Ser Lys Thr Thr Glu Thr Phe Arg Pro Gly
 410 415 420
 Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr
 425 430 435
 30 Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala Pro Thr Arg Ala
 440 445 450
 Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala Val Gly Ile Gly
 455 460 465
 35 Ala Val Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa
 470 475 478

- 40 (2) INFORMATION FOR SEQ ID NO:27:
 (i) SEQUENCE CHARACTERISTICS:
 (A) LENGTH: 1435 base pairs
 (B) TYPE: Nucleic Acid
 (C) STRANDEDNESS: Single
 (D) TOPOLOGY: Linear
 45 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:27:

CTC GAG GTA CCT GTG TGG AAA GAA GCA ACC ACC ACT 36
 Leu Glu Val Pro Val Trp Lys Glu Ala Thr Thr Thr
 1 5 10
 50 CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT GAT TCA GAG 75
 Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Ser Glu
 15 20 25
 55 GCA CAT AAT GTT TGC GCC ACA CAT GCC TGT GTA CCC ACA 114
 Ala His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr
 30 35
 60 GAC CCC AAC CCA CAA GAA GTA GAA TTG GAA AAT GTG ACA 153
 Asp Pro Asn Pro Gln Glu Val Glu Leu Glu Asn Val Thr
 40 45 50

| | | |
|----|---|-----|
| | GAA AAT TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAG | 192 |
| | Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln | |
| | 55 60 | |
| 5 | ATG CAT GGG GAT ATA ATT AGT TTA TGG GAT CAA AGC CTA | 231 |
| | Met His Gly Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu | |
| | 65 70 75 | |
| 10 | AAG CCA TGT GTA AAA TTA ACC CCA CTC TGT GTT ACG TTA | 270 |
| | Lys Pro Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu | |
| | 80 85 90 | |
| 15 | AAT TGC ACT CAC CCA AAT GTT ACT AAT ACC GAG AGA ACG | 309 |
| | Asn Cys Thr Asp Pro Asn Val Thr Asn Ser Glu Arg Thr | |
| | 95 100 | |
| 20 | ATA GAG GGG GGA GAA ATA AAA AAT TGC TCT TTC AAT ATC | 348 |
| | Ile Glu Gly Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile | |
| | 105 110 115 | |
| | ACC ACA AAC ATA AGA GAT AGG TTT CAG AAA GAA TAT GCA | 387 |
| | Thr Thr Asn Ile Arg Asp Arg Phe Gln Lys Glu Tyr Ala | |
| | 120 125 | |
| 25 | CTT TTT TAT AAA CTT GAT GTA ATA CCA TTA GGT AAT GAT | 426 |
| | Leu Phe Tyr Lys Leu Asp Val Ile Pro Leu Gly Asn Asp | |
| | 130 135 140 | |
| 30 | AAT ACT AGC TAT AGG TTG ATA AGT TGT AAC ACC TCA GTC | 465 |
| | Asn Thr Ser Tyr Arg Leu Ile Ser Cys Asn Thr Ser Val | |
| | 145 150 155 | |
| 35 | ATT ACA CAG GCC TGT CCA AAG GTA TCC TTT GAG CCA ATT | 504 |
| | Ile Thr Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile | |
| | 160 165 | |
| 40 | CCC ATA CAT TAT TGT GCC CCG GCT GGT TTT GCG ATT CTA | 543 |
| | Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile Leu | |
| | 170 175 180 | |
| | AAG TGT AAA GAT AAG AAG TTC AAT GGA ACA GGA CCA TGT | 582 |
| | Lys Cys Lys Asp Lys Lys Phe Asn Gly Thr Gly Pro Cys | |
| | 185 190 | |
| 45 | ACA AAT GTC AGC ACA GTA CAA TGT ACA CAT GGA ATT AAG | 621 |
| | Thr Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Lys | |
| | 195 200 205 | |
| 50 | CCA GTA GTA TCA ACT CAA CTG TTG TTA AAT GGC AGT CTA | 660 |
| | Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu | |
| | 210 215 220 | |
| 55 | GCA GAA GAA GAC ATA GTA ATT AGA TCC GCC AAT CTC ACA | 699 |
| | Ala Glu Glu Asp Ile Val Ile Arg Ser Ala Asn Leu Thr | |
| | 225 230 | |
| | GAC AAT GCT AAA AAC ATA ATA GTA CAG CTG AAT GAA TCT | 738 |
| | Asp Asn Ala Lys Asn Ile Ile Val Gln Leu Asn Glu Ser | |
| | 235 240 245 | |
| 60 | GTA ACA ATG AAT TGT ACA AGA CCC AAC AAC AAT ACA ATG | 777 |
| | Val Thr Met Asn Cys Thr Arg Pro Asn Asn Asn Thr Met | |
| | 250 255 | |

AAA AGT ATA CAT ATA GGA CCA GGC AGA GCA TTT TAT GCA 816
 Lys Ser Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala
 260 265 270

5 ACA GGA AAC ATA ATA GGA GAT ATA AGA CAA GCA CAT TGT 855
 Thr Gly Asn Ile Ile Gly Asp Ile Arg Gln Ala His Cys
 275 280 285

10 AAC ATT AGT GGA ACA AAA TGG AAT GAC ACT TTG AAA AAG 894
 Asn Ile Ser Gly Thr Lys Trp Asn Asp Thr Leu Lys Lys
 290 295

15 ATA GCT ATA AAA TTA AGA GAA CAA TTT AAT AAG ACA ATA 933
 Ile Ala Ile Lys Leu Arg Glu Gln Phe Asn Lys Thr Ile
 300 305 310

20 GTC TTT AAT CAA TCC TCA GGA GGG GAC CCA GAA ATT GCA 972
 Val Phe Asn Gln Ser Ser Gly Gly Asp Pro Glu Ile Ala
 315 320

25 ACG CTC AGT TTT AAT TGT GGA GGG GAA TTT TTC TAC TGT 1011
 Thr Leu Ser Phe Asn Cys Gly Gly Glu Phe Phe Tyr Cys
 325 330 335

30 AAT TCA ACA CAA CTG TTT AAT AGT ACT TGG AAT AGT ACT 1050
 Asn Ser Thr Gln Leu Phe Asn Ser Thr Trp Asn Ser Thr
 340 345 350

35 GGG TCA AAT AAC ACT AAA GGA AAT GAC ACA ATC ACA CTC 1089
 Gly Ser Asn Asn Thr Lys Gly Asn Asp Thr Ile Thr Leu
 355 360

40 CCA TGC AGA ATA AGA CAA ATT ATA AAC ATG TGG CAG AAA 1128
 Pro Cys Arg Ile Arg Gln Ile Ile Asn Met Trp Gln Lys
 365 370 375

45 ATA GGA AAA GCA ATG TAT GCC CCT CCC ATC AAA GGG CAA 1167
 Ile Gly Lys Ala Met Tyr Ala Pro Pro Ile Lys Gly Gln
 380 385

50 ATT AGA TGT TCA TCA AAT ATT ACA GGG CTA ATA TTA ACA 1206
 Ile Arg Cys Ser Ser Asn Ile Thr Gly Leu Ile Leu Thr
 390 395 400

55 AGA CAT GGT GGT AAC AAC AAC ATG AGC AAG ACC ACC GAG 1245
 Arg Asp Gly Gly Asn Asn Asn Met Ser Lys Thr Thr Glu
 405 410 415

60 ACC TTC AGA CCT GGA GGA GGA GAT ATG AGG GAC AAT TGG 1284
 Thr Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn Trp
 420 425

65 AGA AGT GAA TTA TAT AAA TAT AAA GTA GTA AAA ATT GAA 1323
 Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu
 430 435 440

70 CCA TTA GGA GTA GCA CCC ACC AGG GCA AAG AGA AGA GTG 1362
 Pro Leu Gly Val Ala Pro Thr Arg Ala Lys Arg Arg Val
 445 450

75 GTG CAG AGA GAA AAA AGA GCA GTG GGA ATA GGA GCT GTG 1401
 Val Gln Arg Glu Lys Arg Ala Val Gly Ile Gly Ala Val
 455 460 465

TTC CTT GGG TTC TTG GGA GCA TAA AGC TTC TAG A 1435
 Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa
 470 475 478

5 (2) INFORMATION FOR SEQ ID NO:28:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 478 amino acids

(B) TYPE: Amino Acid

(D) TOPOLOGY: Linear

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:28:

| | | | | | | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|----|
| | Leu | Glu | Val | Pro | Val | Trp | Lys | Glu | Ala | Thr | Thr | Thr | Leu | Phe | Cys | 1 | 5 | 10 | 15 |
| 15 | Ala | Ser | Asp | Ala | Lys | Ala | Tyr | Asp | Ser | Glu | Ala | His | Asn | Val | Trp | 20 | 25 | 30 | |
| | Ala | Thr | His | Ala | Cys | Val | Pro | Thr | Asp | Pro | Asn | Pro | Gln | Glu | Val | 35 | 40 | 45 | |
| 20 | Glu | Leu | Glu | Asn | Val | Thr | Glu | Asn | Phe | Asn | Met | Trp | Lys | Asn | Asn | 50 | 55 | 60 | |
| | Met | Val | Glu | Gln | Met | His | Gly | Asp | Ile | Ile | Ser | Leu | Trp | Asp | Gln | 65 | 70 | 75 | |
| 25 | Ser | Leu | Lys | Pro | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Val | Thr | Leu | 80 | 85 | 90 | |
| | Asn | Cys | Thr | Asp | Pro | Asn | Val | Thr | Asn | Ser | Glu | Arg | Thr | Ile | Glu | 95 | 100 | 105 | |
| 30 | Gly | Gly | Glu | Ile | Lys | Asn | Cys | Ser | Phe | Asn | Ile | Thr | Thr | Asn | Ile | 110 | 115 | 120 | |
| 35 | Arg | Asp | Arg | Phe | Gln | Lys | Glu | Tyr | Ala | Leu | Phe | Tyr | Lys | Leu | Asp | 125 | 130 | 135 | |
| | Val | Ile | Pro | Leu | Gly | Asn | Asp | Asn | Thr | Ser | Tyr | Arg | Leu | Ile | Ser | 140 | 145 | 150 | |
| 40 | Cys | Asn | Thr | Ser | Val | Ile | Thr | Gln | Ala | Cys | Pro | Lys | Val | Ser | Phe | 155 | 160 | 165 | |
| | Glu | Pro | Ile | Pro | Ile | His | Tyr | Cys | Ala | Pro | Ala | Gly | Phe | Ala | Ile | 170 | 175 | 180 | |
| 45 | Leu | Lys | Cys | Lys | Asp | Lys | Lys | Phe | Asn | Gly | Thr | Gly | Pro | Cys | Thr | 185 | 190 | 195 | |
| 50 | Asn | Val | Ser | Thr | Val | Gln | Cys | Thr | His | Gly | Ile | Lys | Pro | Val | Val | 200 | 205 | 210 | |
| | Ser | Thr | Gln | Leu | Leu | Leu | Asn | Gly | Ser | Leu | Ala | Glu | Glu | Asp | Ile | 215 | 220 | 225 | |
| 55 | Val | Ile | Arg | Ser | Ala | Asn | Leu | Thr | Asp | Asn | Ala | Lys | Asn | Ile | Ile | 230 | 235 | 240 | |
| | Val | Gln | Leu | Asn | Glu | Ser | Val | Thr | Met | Asn | Cys | Thr | Arg | Pro | Asn | 245 | 250 | 255 | |
| 60 | Asn | Asn | Thr | Met | Lys | Ser | Ile | His | Ile | Gly | Pro | Gly | Arg | Ala | Phe | 260 | 265 | 270 | |
| 65 | | | | | | | | | | | | | | | | | | | |

Tyr Ala Thr Gly Asn Ile Ile Gly Asp Ile Arg Gln Ala His Cys
 275 280 285
 5 Asn Ile Ser Gly Thr Lys Trp Asn Asp Thr Leu Lys Lys Ile Ala
 290 295 300
 Ile Lys Leu Arg Glu Gln Phe Asn Lys Thr Ile Val Phe Asn Gln
 305 310 315
 10 Ser Ser Gly Gly Asp Pro Glu Ile Ala Thr Leu Ser Phe Asn Cys
 320 325 330
 Gly Gly Glu Phe Phe Tyr Cys Asn Ser Thr Gln Leu Phe Asn Ser
 335 340 345
 15 Thr Trp Asn Ser Thr Gly Ser Asn Asn Thr Lys Gly Asn Asp Thr
 350 355 360
 Ile Thr Leu Pro Cys Arg Ile Arg Gln Ile Ile Asn Met Trp Gln
 365 370 375
 20 Lys Ile Gly Lys Ala Met Tyr Ala Pro Pro Ile Lys Gly Gln Ile
 380 385 390
 25 Arg Cys Ser Ser Asn Ile Thr Gly Leu Ile Leu Thr Arg Asp Gly
 395 400 405
 Gly Asn Asn Asn Met Ser Lys Thr Thr Glu Thr Phe Arg Pro Gly
 410 415 420
 30 Gly Gly Asp Met Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr
 425 430 435
 35 Lys Val Val Lys Ile Glu Pro Leu Gly Val Ala Pro Thr Arg Ala
 440 445 450
 Lys Arg Arg Val Val Gln Arg Glu Lys Arg Ala Val Gly Ile Gly
 455 460 465
 40 Ala Val Phe Leu Gly Phe Leu Gly Ala Xaa Ser Phe Xaa
 470 475 478

(2) INFORMATION FOR SEQ ID NO:29:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 511 amino acids

(B) TYPE: Amino Acid

(D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:29:

50 Met Arg Val Lys Gly Ile Arg Arg Asn Tyr Gln His Trp Trp Gly Arg
 1 5 10 15
 Gly Thr Met Leu Leu Gly Leu Leu Met Ile Cys Ser Ala Thr Glu Lys
 20 25 30
 55 Leu Trp Val Thr Val Tyr Tyr Gly Val Pro Val Trp Lys Glu Ala Thr
 35 40 45
 Thr Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu Ala
 50 55 60
 60 His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp Pro Asn Pro
 65 70 75 80

| | | | | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|--|
| | Gln | Glu | Val | Glu | Leu | Val | Asn | Val | Thr | Glu | Asn | Phe | Asn | Met | Trp | Lys | |
| | | | | | 85 | | | | | 90 | | | | | 95 | | |
| 5 | Asn | Asn | Met | Val | Glu | Gln | Met | His | Glu | Asp | Ile | Ile | Ser | Leu | Trp | Asn | |
| | | | | 100 | | | | | 105 | | | | | 110 | | | |
| | Gln | Ser | Leu | Lys | Pro | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Val | Thr | Leu | |
| | | | | 115 | | | | 120 | | | | | 125 | | | | |
| 10 | Asn | Cys | Thr | Asp | Leu | Arg | Asn | Thr | Thr | Asn | Thr | Asn | Asn | Ser | Thr | Asp | |
| | | 130 | | | | | 135 | | | | | 140 | | | | | |
| | Asn | Asn | Asn | Ser | Lys | Ser | Glu | Gly | Thr | Ile | Lys | Gly | Gly | Glu | Met | Lys | |
| | 145 | | | | | 150 | | | | | 155 | | | | | 160 | |
| 15 | Asn | Cys | Ser | Phe | Asn | Ile | Thr | Thr | Ser | Ile | Gly | Asp | Lys | Met | Gln | Lys | |
| | | | | 165 | | | | | | 170 | | | | | 175 | | |
| 20 | Glu | Tyr | Ala | Leu | Leu | Tyr | Lys | Leu | Asp | Ile | Glu | Pro | Ile | Asp | Asn | Asp | |
| | | | | 180 | | | | | 185 | | | | | 190 | | | |
| | Ser | Thr | Ser | Tyr | Arg | Leu | Ile | Ser | Cys | Asn | Thr | Ser | Val | Ile | Thr | Gln | |
| | | | 195 | | | | | 200 | | | | | 205 | | | | |
| 25 | Ala | Cys | Pro | Lys | Ile | Ser | Phe | Glu | Pro | Ile | Pro | Ile | His | Tyr | Cys | Ala | |
| | | 210 | | | | | 215 | | | | | 220 | | | | | |
| | Pro | Ala | Gly | Phe | Ala | Ile | Leu | Lys | Cys | Asn | Asp | Lys | Lys | Phe | Ser | Gly | |
| | 225 | | | | 230 | | | | | | 235 | | | | | 240 | |
| 30 | Lys | Gly | Ser | Cys | Lys | Asn | Val | Ser | Thr | Val | Gln | Cys | Thr | His | Gly | Ile | |
| | | | | | 245 | | | | | 250 | | | | | 255 | | |
| 35 | Arg | Pro | Val | Val | Ser | Thr | Gln | Leu | Leu | Leu | Asn | Gly | Ser | Leu | Ala | Glu | |
| | | | 260 | | | | | | 265 | | | | | 270 | | | |
| | Glu | Glu | Val | Val | Ile | Arg | Ser | Glu | Asp | Phe | Thr | Asp | Asn | Ala | Lys | Thr | |
| | | | 275 | | | | | 280 | | | | | 285 | | | | |
| 40 | Ile | Ile | Val | His | Leu | Lys | Glu | Ser | Val | Gln | Ile | Asn | Cys | Thr | Arg | Pro | |
| | | 290 | | | | 295 | | | | | | 300 | | | | | |
| | Asn | Tyr | Asn | Lys | Arg | Lys | Arg | Ile | His | Ile | Gly | Pro | Gly | Arg | Ala | Phe | |
| | 305 | | | | 310 | | | | | | 315 | | | | | 320 | |
| 45 | Tyr | Thr | Thr | Lys | Asn | Ile | Lys | Gly | Thr | Ile | Arg | Gln | Ala | His | Cys | Ile | |
| | | | | 325 | | | | | | 330 | | | | | 335 | | |
| 50 | Ile | Ser | Arg | Ala | Lys | Trp | Asn | Asp | Thr | Leu | Arg | Gln | Ile | Val | Ser | Lys | |
| | | | | 340 | | | | | 345 | | | | | 350 | | | |
| | Leu | Lys | Glu | Gln | Phe | Lys | Asn | Lys | Thr | Ile | Val | Phe | Asn | Pro | Ser | Ser | |
| | | | 355 | | | | | 360 | | | | | 365 | | | | |
| 55 | Gly | Gly | Asp | Pro | Glu | Ile | Val | Met | His | Ser | Phe | Asn | Cys | Gly | Gly | Glu | |
| | | 370 | | | | 375 | | | | | | 380 | | | | | |
| | Phe | Phe | Tyr | Cys | Asn | Thr | Ser | Pro | Leu | Phe | Asn | Ser | Ile | Trp | Asn | Gly | |
| | 385 | | | | 390 | | | | | | 395 | | | | | 400 | |
| 60 | Asn | Asn | Thr | Trp | Asn | Asn | Thr | Thr | Gly | Ser | Asn | Asn | Asn | Ile | Thr | Leu | |
| | | | | 405 | | | | | | 410 | | | | 415 | | | |
| 65 | Gln | Cys | Lys | Ile | Lys | Gln | Ile | Ile | Asn | Met | Trp | Gln | Lys | Val | Gly | Lys | |
| | | | 420 | | | | | 425 | | | | | 430 | | | | |

Ala Met Tyr Ala Pro Pro Ile Glu Gly Gln Ile Arg Cys Ser Ser Asn
 435 440 445
 5 Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Glu Asp Thr Asp Thr
 450 455 460
 Asn Asp Thr Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn
 465 470 475 480
 10 Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Thr Ile Glu Pro Leu
 485 490 495
 Gly Val Ala Pro Thr Lys Ala Lys Arg Arg Val Val Gln Arg Glu
 500 505 510
 15

(2) INFORMATION FOR SEQ ID NO:30:

(i) SEQUENCE CHARACTERISTICS:

- (A) LENGTH: 2800 base pairs
 (B) TYPE: Nucleic Acid
 (C) STRANDEDNESS: Single
 (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:30:

25 TTCGAGCTCG CCCGACATTG ATTATTGACT AGAGTCGATC GACAGCTGTG 50
 GAATGTGTGT CAGTTAGGGT GTGGAAAGTC CCCAGGCTCC CCAGCAGGCA 100
 GAAGTATGCA AAGCATGCAT CTCAATTAGT CAGCAACCAG GTGTGGAAAG 150
 30 TCCCCAGGCT CCCCAGCAGG CAGAAGTATG CAAAGCATGC ATCTCAATTA 200
 GTCAGCAACC ATAGTCCCGC CCCTAACTEC GCCCATCCCG CCCCTAACTC 250
 35 CGCCCAGTTC CGCCCATTCT CCGCCCCATG GCTGACTAAT TTTTTTTATT 300
 TATGCAGAGG CCGAGGCCCG CTCGGCCTCT GAGCTATTCC AGAAGTAGTG 350
 AGGAGGCTTT TTTGGAGGCC TAGGCTTTTG CAAAAGCTA GCTTATCCGG 400
 40 CCGGGAACGG TGCATTGGAA CGCGGATTCC CCGTGCCAAG ACTCAGGTAA 450
 GTACCGCCTA TAGAGTCTAT AGGCCCCACC CTTGGCTTC GTTAGAACGC 500
 45 GGCTACAATT AATACATAAC CTTTTGGATC GATCCTACTG ACACTGACAT 550
 CCACTTTTTC TTTTCTCCA CAGGTGTCCA CTCCCAGGTC CAACTGCACC 600
 TCGGTTTCGG AAGCTAGCTT GGGCTGCATC GATTGAATTC CACTGCCTTC 650
 50 CACCAAGCTC TGCAGGATCC CAGAGTCAGG GG TCT GTA TCT TCC TGC 697
 Ser Val Ser Ser Cys
 1 5
 55 TGG TGG CTC CAG TTC AGG AAC AGT AAA CCC TGC TCC GAA TAT 739
 Trp Trp Leu Gln Phe Arg Asn Ser Lys Pro Cys Ser Glu Tyr
 10 15
 60 TGC CTC TCA CAT CTC GTC AAT CTC CGC GAG GAC TGG GGA CCC 781
 Cys Leu Ser His Leu Val Asn Leu Arg Glu Asp Trp Gly Pro
 20 25 30

| | | | | | | | | | | | | | | | |
|----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|-----|------|
| | TCT | GAC | AAG | CTT | CAG | CGC | GAA | CGA | CCA | ACT | ACC | CCG | ATC | ATC | 823 |
| | Cys | Asp | Lys | Leu | Gln | Arg | Glu | Arg | Pro | Thr | Thr | Pro | Ile | Ile | |
| | 35 | | | | | | 40 | | | | | 45 | | | |
| 5 | AGT | TAT | CCT | TAA | GGT | CTC | TTT | TGT | GTG | GTG | CGT | TCC | GGT | ATG | 865 |
| | Ser | Tyr | Pro | * | Gly | Leu | Phe | Cys | Val | Val | Arg | Ser | Gly | Met | |
| | 50 | | | | | | 55 | | | | | | 60 | | |
| 10 | GGG | GGG | ACT | GCC | GCC | AGG | TTG | GGG | GCC | GTG | ATT | TTG | TTT | GTC | 907 |
| | Gly | Gly | Thr | Ala | Ala | Arg | Leu | Gly | Ala | Val | Ile | Leu | Phe | Val | |
| | 62 | | | 65 | | | | 70 | | | | | 75 | | |
| 15 | GTC | ATA | GTG | GGC | CTC | CAT | GGG | GTC | CGC | GGC | AAA | TAT | GCC | TTG | 949 |
| | Val | Ile | Val | Gly | Leu | His | Gly | Val | Arg | Gly | Lys | Tyr | Ala | Leu | |
| | | | | 80 | | | | | | 85 | | | | | |
| 20 | GCG | GAT | GCC | TCT | CTC | AAG | ATG | GCC | GAC | CCC | AAT | CGA | TTT | CGC | 991 |
| | Ala | Asp | Ala | Ser | Leu | Lys | Met | Ala | Asp | Pro | Asn | Arg | Phe | Arg | |
| | 90 | | | | | 95 | | | | | 100 | | | | |
| 25 | GGC | AAA | GAC | CTT | CCG | GTC | CTG | GAC | CAG | CTG | CTC | GAG | GTA | CCT | 1033 |
| | Gly | Lys | Asp | Leu | Pro | Val | Leu | Asp | Gln | Leu | Leu | Glu | Val | Pro | |
| | | | 105 | | | | | 110 | | | | | 115 | | |
| | GTG | TGG | AAA | GAA | GCA | AAC | ACC | ACT | CTA | TTT | TGT | GCA | TCA | GAT | 1075 |
| | Val | Trp | Lys | Glu | Ala | Asn | Thr | Thr | Leu | Phe | Cys | Ala | Ser | Asp | |
| | | | | 120 | | | | 125 | | | | | 130 | | |
| 30 | GCT | AAA | GCA | TAT | AAG | ACA | GAG | GCA | CAT | AAT | GTT | TGG | GCC | ACA | 1117 |
| | Ala | Lys | Ala | Tyr | Lys | Thr | Glu | Ala | His | Asn | Val | Trp | Ala | Thr | |
| | | | | 135 | | | | 140 | | | | | 145 | | |
| 35 | CAT | GCC | TGT | GTA | CCC | ACA | GAC | CCC | AAA | CCA | CAA | GAA | ATA | AAA | 1159 |
| | His | Ala | Cys | Val | Pro | Thr | Asp | Pro | Lys | Pro | Gln | Glu | Ile | Lys | |
| | | | | | 150 | | | | | 155 | | | | | |
| 40 | TTG | GAA | AAT | GTG | ACA | GAA | AAT | TTT | AAC | ATG | TGG | AAA | AAT | AAC | 1201 |
| | Leu | Glu | Asn | Val | Thr | Glu | Asn | Phe | Asn | Met | Trp | Lys | Asn | Asn | |
| | 160 | | | | | 165 | | | | | 170 | | | | |
| 45 | ATG | GTA | GAA | CAG | ATG | CAT | GAG | GAT | ATA | ATC | AGT | TTA | TGG | GAT | 1243 |
| | Met | Val | Glu | Gln | Met | His | Glu | Asp | Ile | Ile | Ser | Leu | Trp | Asp | |
| | | 175 | | | | | 180 | | | | | 185 | | | |
| | CAA | AGC | CTA | AAG | CCA | TGT | GTA | AAA | TTA | ACC | CCA | CTC | TGT | GTT | 1285 |
| | Gln | Ser | Leu | Lys | Pro | Cys | Val | Lys | Leu | Thr | Pro | Leu | Cys | Val | |
| | | | 190 | | | | | 195 | | | | | 200 | | |
| 50 | ACT | TTA | AAT | TGC | ACT | GAT | TTG | AGG | AAT | AAT | ACT | AAT | ACC | AAT | 1327 |
| | Thr | Leu | Asn | Cys | Thr | Asp | Leu | Arg | Asn | Asn | Thr | Asn | Thr | Asn | |
| | | | | 205 | | | | 210 | | | | | 215 | | |
| 55 | AGT | ACC | TAC | GGA | AAA | ATA | ATG | GAG | GGA | GGA | GAG | ATA | AAA | AAC | 1369 |
| | Ser | Thr | Tyr | Gly | Lys | Ile | Met | Glu | Gly | Gly | Glu | Ile | Lys | Asn | |
| | | | | 220 | | | | | 225 | | | | | | |
| 60 | TGC | TCT | TTC | AAT | ATC | ACC | ACA | AGC | ATA | AAA | GAT | AAG | CTG | AAA | 1411 |
| | Cys | Ser | Phe | Asn | Ile | Thr | Thr | Ser | Ile | Lys | Asp | Lys | Leu | Lys | |
| | 230 | | | | | 235 | | | | | 240 | | | | |
| 65 | GAT | ATG | TCA | CTT | TTT | TAT | AAA | CTT | GAT | GTA | GTA | CCA | ATA | GGT | 1453 |
| | Asp | Met | Ser | Leu | Phe | Tyr | Lys | Leu | Asp | Val | Val | Pro | Ile | Gly | |
| | 245 | | | | | | 250 | | | | | 255 | | | |

| | | |
|----|---|------|
| | AAT AAT AGT AAT ACT ACT AGT TAT AGG TTG ATA AGT TGT AAC | 1495 |
| | Asn Asn Ser Asn Thr Thr Ser Tyr Arg Leu Ile Ser Cys Asn | |
| | 260 265 270 | |
| 5 | ACC TCA GTC ATT ACA CAA GCC TGT CCA AAG ACA TCC TTT GAG | 1537 |
| | Thr Ser Val Ile Thr Gln Ala Cys Pro Lys Thr Ser Phe Glu | |
| | 275 280 285 | |
| 10 | CCA ATT CCC ATA CAT TAT TGT GCC CCG GCT GGT TTT GCG ATT | 1579 |
| | Pro Ile Pro Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile | |
| | 290 295 | |
| 15 | CTC AAG TGT AAT GAT AAT AAG TTC AAT GGA ACA GGA CCA TGT | 1621 |
| | Leu Lys Cys Asn Asp Asn Lys Phe Asn Gly Thr Gly Pro Cys | |
| | 300 305 310 | |
| 20 | CCA AAT GTC AGC ACA GTA CAA TGT ACA CAT GGA ATT AGG CCA | 1663 |
| | Pro Asn Val Ser Thr Val Gln Cys Thr His Gly Ile Arg Pro | |
| | 315 320 325 | |
| | GTA GTA TCA ACT CAA CTG CTG TTA AAT GGC AGT CTA GCA GAA | 1705 |
| | Val Val Ser Thr Gln Leu Leu Asn Gly Ser Leu Ala Glu | |
| | 330 335 340 | |
| 25 | AAA GAG GTA GTC CTT AGA TCT GAA AAT TTC ACG GAC AAT GCT | 1747 |
| | Lys Glu Val Val Leu Arg Ser Glu Asn Phe Thr Asp Asn Ala | |
| | 345 350 355 | |
| 30 | AAA ACC ATA ATA GTA CAG CTG AAC GAA TCT GTA ATA ATT GAT | 1789 |
| | Lys Thr Ile Ile Val Gln Leu Asn Glu Ser Val Ile Ile Asp | |
| | 360 365 | |
| 35 | TGT ATG AGA CCC AAC AAC AAT ACA AGA ACA AGT ATA CCT ATG | 1831 |
| | Cys Met Arg Pro Asn Asn Asn Thr Arg Thr Ser Ile Pro Met | |
| | 370 375 380 | |
| 40 | GGA CCA GGG AAA GCA TTT TAT GCA ACA GGA GAT GTA ATA GGA | 1873 |
| | Gly Pro Gly Lys Ala Phe Tyr Ala Thr Gly Asp Val Ile Gly | |
| | 385 390 395 | |
| | GAT ATA AGA CGA GCA CAT TGT AAC ATT AGT AGA GCA GGA TGG | 1915 |
| | Asp Ile Arg Arg Ala His Cys Asn Ile Ser Arg Ala Gly Trp | |
| | 400 405 410 | |
| 45 | AAT ACC ACT TTA CAA CAG ATA GCT AAA AAA TTA AGA GAA AAA | 1957 |
| | Asn Thr Thr Leu Gln Gln Ile Ala Lys Lys Leu Arg Glu Lys | |
| | 415 420 425 | |
| 50 | TTT GAG AAC AAA ACA ATA GTT TTT AAT CAC TCC TCA GCA GGG | 1999 |
| | Phe Glu Asn Lys Thr Ile Val Phe Asn His Ser Ser Gly Gly | |
| | 430 435 | |
| 55 | GAC CCA GAA ATT GTA ATG CAC ACT TTT AAT TGT GGA GGG GAA | 2041 |
| | Asp Pro Glu Ile Val Met His Thr Phe Asn Cys Gly Gly Glu | |
| | 440 445 450 | |
| 60 | TTT TTC TGC TGT AAT TCA ACA CCA CTG TTT AAT AGT ACT TGG | 2083 |
| | Phe Phe Cys Cys Asn Ser Thr Pro Leu Phe Asn Ser Thr Trp | |
| | 455 460 465 | |
| | AAT GAT GCA CAA CTG TTT AAT AGT ACT TGG GAT GAT ACT AAA | 2125 |
| | Asn Asp Ala Gln Leu Phe Asn Ser Thr Trp Asp Asp Thr Lys | |
| | 470 475 480 | |
| 65 | | |

TGG TCA AAA GGC ACT AAC GAA AAT GAC ACA ATC ACC CTC CAT 2167
 Trp Ser Lys Gly Thr Asn Glu Asn Asp Thr Ile Thr Leu His
 485 490 495
 5 TGC AGA ATA AAA CAA ATT ATA AAT ATG TGG CAG GAA GTA GGA 2209
 Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly
 500 505
 10 AAA GCA ATG TAT GCC CCT CCC ATC AAA GGA CAA ATT AGA TGT 2251
 Lys Ala Met Tyr Ala Pro Pro Ile Lys Gly Gln Ile Arg Cys
 510 515 520
 GAA TCA AAT ATT ACA GGG CTG CTA TTA ACA AGA GAT GGT GGT 2293
 15 Glu Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly
 525 530 535
 20 AAC GAC ACG AGC AAG AAT AAC ACT GAG ATT TTC AGA CCT GGA 2335
 Asn Asp Thr Ser Lys Asn Asn Thr Glu Ile Phe Arg Pro Gly
 540 545 550
 GGA GGA AAT ATG AAG GAC AAT TGG AGA ACT GAA TTA TAT AAA 2377
 Gly Gly Asn Met Lys Asp Asn Trp Arg Ser Glu Leu Tyr Lys
 555 560 565
 25 TAT AAA GTA ATA AAA ATT GAA CCA TTA GGA GTA GCA CCC ATC 2419
 Tyr Lys Val Ile Lys Ile Glu Pro Leu Gly Val Ala Pro Ile
 570 575 579
 30 TAGGCAAAGA GAAGAGTGGT GCAGAGAGAA AAAAGAGCAG TGACACTAGG 2469
 35 AGCTATGTTC CTTGGGTTCT TGGGAGCAGC AGGAAGCACT ATGGGCGATA 2519
 AGCTTTAATG CCGTAGTTTA TCACAGTTAA ATTCGTAACG CACTCAGGCA 2569
 CCGTGTATGA AATCTAACAA TGCGACCTGC AGAAGCTTAG AACCGAGGAA 2619
 40 CTTGTTTATT GCAGCTTATA ATGGTTACAA ATAAAGCAAT ACCATCACAA 2669
 ATTTACACAA TAAAGCATTT TTTTCACTGC ATTCTAGTTG TGGTTTGTCC 2719
 45 AAACATCATCA ATGTATCTTA TCATGTCTGG ATCGGGAATT AATTGGGCGC 2769
 AGCACCATGG CCTGAAATAA CCTCTGAAAG A 2800

50 (2) INFORMATION FOR SEQ ID NO:31

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 579 amino acids

(B) TYPE: Amino Acid

(D) TOPOLOGY: Linear

55 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:31:

Ser Val Ser Ser Cys Trp Trp Leu Gln Phe Arg Asn Ser Lys Pro Cys
 1 5 10 15
 60 Ser Glu Tyr Cys Leu Ser His Leu Val Asn Leu Arg Glu Asp Trp Gly
 20 25 30
 Pro Cys Asp Lys Leu Gln Arg Glu Arg Pro Thr Thr Pro Ile Ile Ser
 35 40 45
 65

Tyr Pro * Gly Leu Phe Cys Val Val Arg Ser Gly Met Gly Gly Thr
 50 55 60
 5 Ala Ala Arg Leu Gly Ala Val Ile Leu Phe Val Val Ile Val Gly Leu
 65 70 75 80
 His Gly Val Arg Gly Lys Tyr Ala Leu Ala Asp Ala Ser Leu Lys Met
 85 90 95
 10 Ala Asp Pro Asn Arg Phe Arg Gly Lys Asp Leu Pro Val Leu Asp Gln
 100 105 110
 Leu Leu Glu Val Pro Val Trp Lys Glu Ala Asn Thr Thr Leu Phe Cys
 115 120 125
 15 Ala Ser Asp Ala Lys Ala Tyr Lys Thr Glu Ala His Asn Val Trp Ala
 130 135 140
 Thr His Ala Cys Val Pro Thr Asp Pro Lys Pro Gln Glu Ile Lys Leu
 145 150 155 160
 Glu Asn Val Thr Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu
 165 170 175
 25 Gln Met His Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys Pro
 180 185 190
 Cys Val Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys Thr Asp Leu
 195 200 205
 30 Arg Asn Asn Thr Asn Thr Asn Ser Thr Tyr Gly Lys Ile Met Glu Gly
 210 215 220
 Gly Glu Ile Lys Asn Cys Ser Phe Asn Ile Thr Thr Ser Ile Lys Asp
 225 230 235 240
 Lys Leu Lys Asp Met Ser Leu Phe Tyr Lys Leu Asp Val Val Pro Ile
 245 250 255
 40 Gly Asn Asn Ser Asn Thr Thr Ser Tyr Arg Leu Ile Ser Cys Asn Thr
 260 265 270
 Ser Val Ile Thr Gln Ala Cys Pro Lys Thr Ser Phe Glu Pro Ile Pro
 275 280 285
 45 Ile His Tyr Cys Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Asn Asp
 290 295 300
 Asn Lys Phe Asn Gly Thr Gly Pro Cys Pro Asn Val Ser Thr Val Gln
 305 310 315 320
 Cys Thr His Gly Ile Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn
 325 330 335
 55 Gly Ser Leu Ala Glu Lys Glu Val Val Leu Arg Ser Glu Asn Phe Thr
 340 345 350
 Asp Asn Ala Lys Thr Ile Ile Val Gln Leu Asn Glu Ser Val Ile Ile
 355 360 365
 60 Asp Cys Met Arg Pro Asn Asn Asn Thr Arg Thr Ser Ile Pro Met Gly
 370 375 380
 Pro Gly Lys Ala Phe Tyr Ala Thr Gly Asp Val Ile Gly Asp Ile Arg
 385 390 395 400

Arg Ala His Cys Asn Ile Ser Arg Ala Gly Trp Asn Thr Thr Leu Gln
 405 410 415
 5 Gln Ile Ala Lys Lys Leu Arg Glu Lys Phe Glu Asn Lys Thr Ile Val
 420 425 430
 Phe Asn His Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Thr Phe
 435 440 445
 10 Asn Cys Gly Gly Glu Phe Phe Cys Cys Asn Ser Thr Pro Leu Phe Asn
 450 455 460
 Ser Thr Trp Asn Asp Ala Gln Leu Phe Asn Ser Thr Trp Asp Asp Thr
 465 470 475 480
 15 Lys Trp Ser Lys Gly Thr Asn Glu Asn Asp Thr Ile Thr Leu His Cys
 485 490 495
 20 Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys Ala Met
 500 505 510
 Tyr Ala Pro Pro Ile Lys Gly Gln Ile Arg Cys Glu Ser Asn Ile Thr
 515 520 525
 25 Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Asp Thr Ser Lys Asn Asn
 530 535 540
 Thr Glu Ile Phe Arg Pro Gly Gly Gly Asn Met Lys Asp Asn Trp Arg
 545 550 555 560
 30 Ser Glu Leu Tyr Lys Tyr Lys Val Ile Lys Ile Glu Pro Leu Gly Val
 565 570 575

Ala Pro Ile

(2) INFORMATION FOR SEQ ID NO:32:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1533 base pairs

(B) TYPE: Nucleic Acid

(C) STRANDEDNESS: Single

(D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:32:

45 ATGGGGGGGA CTGCCGCCAG GTTGGGGGCC GTGATTTTGT TTGTCGTCAT 50
 AGTGGGCCTC CATGGGGTCC GCGGCAAATA TGCCTTGGCG GATGCCTCTC 100
 TCAAGATGGC CGACCCCAAT CGATTTGCGG GCAAAGACCT TCCGGTCCTG 150
 50 GACCAGCTGC TCGAG GTA CCT GTG TGG AAA GAA GCA ACC ACC 192
 Val Pro Val Trp Lys Glu Ala Thr Thr
 1 5
 55 ACT CTA TTT TGT GCA TCA GAT GCT AAA GCA TAT GAT ACA GAG 234
 Thr Leu Phe Cys Ala Ser Asp Ala Lys Ala Tyr Asp Thr Glu
 10 15 20
 60 GTA CAT AAT GTT TGG GCC ACA CAT GCC TGT GTA CCC ACA GAC 276
 Val His Asn Val Trp Ala Thr His Ala Cys Val Pro Thr Asp
 25 30 35

| | | |
|----|---|-----|
| | CCC AAC CCA CAA GAA ATA GGA TTG GAA AAT GTA ACA GAA AAT | 318 |
| | Pro Asn Pro Gln Glu Ile Gly Leu Glu Asn Val Thr Glu Asn | |
| | 40 45 50 | |
| 5 | TTT AAC ATG TGG AAA AAT AAC ATG GTA GAA CAG ATG CAT GAG | 360 |
| | Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His Glu | |
| | 55 60 65 | |
| 10 | GAT ATA ATC AGT TTA TGG GAT CAA AGC TTA AAG CCA TGT GTA | 402 |
| | Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys Pro Cys Val | |
| | 70 75 | |
| 15 | AAA TTA ACC CCA CTA TGT GTT ACT TTA AAT TGC ACT GAT TTG | 444 |
| | Lys Leu Thr Pro Leu Cys Val Thr Leu Asn Cys Thr Asp Leu | |
| | 80 85 90 | |
| 20 | AAA AAT GCT ACT AAT ACC ACT AGT AGC AGC TGG GGA AAG ATG | 486 |
| | Lys Asn Ala Thr Asn Thr Thr Ser Ser Ser Trp Gly Lys Met | |
| | 95 100 105 | |
| 25 | GAG AGA GGA GAA ATA AAA AAC TGC TCT TTC AAT GTC ACC ACA | 528 |
| | Glu Arg Gly Glu Ile Lys Asn Cys Ser Phe Asn Val Thr Thr | |
| | 110 115 120 | |
| 30 | AGT ATA AGA GAT AAG ATG AAG AAT GAA TAT GCA CTT TTT TAT | 570 |
| | Ser Ile Arg Asp Lys Met Lys Asn Glu Tyr Ala Leu Phe Tyr | |
| | 125 130 135 | |
| 35 | AAA CTT GAT GTA GTA CCA ATA GAT AAT GAT AAT ACT AGC TAT | 612 |
| | Lys Leu Asp Val Val Pro Ile Asp Asn Asp Asn Thr Ser Tyr | |
| | 140 145 | |
| 40 | AGG TTG ATA AGT TGT AAC ACC TCA GTC ATT ACA CAG GCC TGT | 654 |
| | Arg Leu Ile Ser Cys Asn Thr Ser Val Ile Thr Gln Ala Cys | |
| | 150 155 160 | |
| 45 | CCA AAG GTG TCC TTT GAG CCA ATT CCC ATA CAT TAT TGT GCC | 696 |
| | Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys Ala | |
| | 165 170 175 | |
| 50 | CCG GCT GGT TTT GCG ATT CTA AAG TGT AGA GAT AAA AAG TTC | 738 |
| | Pro Ala Gly Phe Ala Ile Leu Lys Cys Arg Asp Lys Lys Phe | |
| | 180 185 190 | |
| 55 | AAC GGA ACA GGA CCA TGT ACA AAT GTC AGC ACA GTA CAA TGT | 780 |
| | Asn Gly Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys | |
| | 195 200 205 | |
| 60 | ACA CAT GGA ATT AGG CCA GTA GTA TCA ACT CAA CTG CTG TTA | 822 |
| | Thr His Gly Ile Arg Pro Val Val Ser Thr Gln Leu Leu Leu | |
| | 210 215 | |
| 65 | AAT GCC AGT TTA GCA GAA GAA GAA GTA GTA ATT ACA TCT GCC | 864 |
| | Asn Gly Ser Leu Ala Glu Glu Glu Val Val Ile Arg Ser Ala | |
| | 220 225 230 | |
| 70 | AAT TTC TCG GAC AAT GCT AAA ACC ATA ATA GTA CAG CTG AAC | 906 |
| | Asn Phe Ser Asp Asn Ala Lys Thr Ile Ile Val Gln Leu Asn | |
| | 235 240 245 | |

| | | |
|----|---|------|
| | GAA TCT GTA GAA ATT AAT TGT ACA AGA CCC AAC AAC AAT ACA | 948 |
| | Glu Ser Val Glu Ile Asn Cys Thr Arg Pro Asn Asn Asn Thr | |
| | 250 255 260 | |
| 5 | AGA AGA ACT ATA CAT ATA GGA CCA GGG AGA GCA TTT TAT GCA | 990 |
| | Arg Arg Ser Ile His Ile Gly Pro Gly Arg Ala Phe Tyr Ala | |
| | 265 270 275 | |
| 10 | ACA GGA GAA ATA ATA GGA GAC ATA AGA CAA GCA CAT TGT AAC | 1032 |
| | Thr Gly Glu Ile Ile Gly Asp Ile Arg Gln Ala His Cys Asn | |
| | 280 285 | |
| 15 | CTT AGT AGC ACA AAA TGG AAT AAT ACT TTA AAA CAG ATA GTT | 1074 |
| | Leu Ser Ser Thr Lys Trp Asn Asn Thr Leu Lys Gln Ile Val | |
| | 290 295 300 | |
| 20 | ACA AAA TTA AGA GAA CAT TTT AAT AAA ACA ATA GTC TTT AAT | 1116 |
| | Thr Lys Leu Arg Glu His Phe Asn Lys Thr Ile Val Phe Asn | |
| | 305 310 315 | |
| | CAC TCC TCA GGA GGG GAC CCA GAA ATT GTA ATG CAC AGT TTT | 1158 |
| | His Ser Ser Gly Gly Asp Pro Glu Ile Val Met His Ser Phe | |
| | 320 325 330 | |
| 25 | AAT TGT GGA GGG GAA TTT TTC TAC TGT AAT ACA ACA CCA CTG | 1200 |
| | Asn Cys Gly Gly Glu Phe Phe Tyr Cys Asn Thr Thr Pro Leu | |
| | 335 340 345 | |
| 30 | TTT AAT AGT ACT TGG AAT TAT ACT TAT ACT TGG AAT AAT ACT | 1242 |
| | Phe Asn Ser Thr Trp Asn Tyr Thr Tyr Thr Trp Asn Asn Thr | |
| | 350 355 | |
| 35 | GAA GGG TCA AAT GAC ACT GGA AGA AAT ATC ACA CTC CAA TGC | 1284 |
| | Glu Gly Ser Asn Asp Thr Gly Arg Asn Ile Thr Leu Gln Cys | |
| | 360 365 370 | |
| 40 | AGA ATA AAA CAA ATT ATA AAC ATG TGG CAG GAA GTA GGA AAA | 1326 |
| | Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu Val Gly Lys | |
| | 375 380 385 | |
| | GCA ATG TAT GCC CCT CCC ATA AGA GGA CAA ATT AGA TGC TCA | 1368 |
| | Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile Arg Cys Ser | |
| | 390 395 400 | |
| 45 | TCA AAT ATT ACA GGG CTG CTA TTA ACA AGA GAT GGT GGT AAT | 1410 |
| | Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn | |
| | 405 410 415 | |
| 50 | AAC AGC GAA ACC GAG ATC TTC AGA CCT GGA GGA GCA GAT ATG | 1452 |
| | Asn Ser Glu Thr Glu Ile Phe Arg Pro Gly Gly Gly Asp Met | |
| | 420 425 | |
| 55 | AGG GAC AAT TGG AGA AGT GAA TTA TAT AAA TAT AAA GTA GTA | 1494 |
| | Arg Asp Asn Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val | |
| | 430 435 440 | |
| 60 | AAA ATT GAA CCA TTA GGA GTA GCA CCC ACC AAG GCA TAA | 1533 |
| | Lys Ile Glu Pro Leu Gly Val Ala Pro Thr Lys Ala * | |
| | 445 450 455 | |

(2) INFORMATION FOR SEQ ID NO:33:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 456 amino acids

(B) TYPE: Amino Acid

(D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:33:

Val Pro Val Trp Lys Glu Ala Thr Thr Thr Leu Phe Cys Ala Ser Asp
 1 5 10 15
 Ala Lys Ala Tyr Asp Thr Glu Val His Asn Val Trp Ala Thr His Ala
 20 25 30
 Cys Val Pro Thr Asp Pro Asn Pro Gln Glu Ile Gly Leu Glu Asn Val
 35 40 45
 Thr Glu Asn Phe Asn Met Trp Lys Asn Asn Met Val Glu Gln Met His
 50 55 60
 Glu Asp Ile Ile Ser Leu Trp Asp Gln Ser Leu Lys Pro Cys Val Lys
 65 70 75 80
 Leu Thr Pro Leu Cys Val Thr Leu Asn Cys Thr Asp Leu Lys Asn Ala
 85 90 95
 Thr Asn Thr Thr Ser Ser Ser Trp Gly Lys Met Glu Arg Gly Glu Ile
 100 105 110
 Lys Asn Cys Ser Phe Asn Val Thr Thr Ser Ile Arg Asp Lys Met Lys
 115 120 125
 Asn Glu Tyr Ala Leu Phe Tyr Lys Leu Asp Val Val Pro Ile Asp Asn
 130 135 140
 Asp Asn Thr Ser Tyr Arg Leu Ile Ser Cys Asn Thr Ser Val Ile Thr
 145 150 155 160
 Gln Ala Cys Pro Lys Val Ser Phe Glu Pro Ile Pro Ile His Tyr Cys
 165 170 175
 Ala Pro Ala Gly Phe Ala Ile Leu Lys Cys Arg Asp Lys Lys Phe Asn
 180 185 190
 Gly Thr Gly Pro Cys Thr Asn Val Ser Thr Val Gln Cys Thr His Gly
 195 200 205
 Ile Arg Pro Val Val Ser Thr Gln Leu Leu Leu Asn Gly Ser Leu Ala
 210 215 220
 Glu Glu Glu Val Val Ile Arg Ser Ala Asn Phe Ser Asp Asn Ala Lys
 225 230 235 240
 Thr Ile Ile Val Gln Leu Asn Glu Ser Val Glu Ile Asn Cys Thr Arg
 245 250 255
 Pro Asn Asn Asn Thr Arg Arg Ser Ile His Ile Gly Pro Gly Arg Ala
 260 265 270
 Phe Tyr Ala Thr Gly Glu Ile Ile Gly Asp Ile Arg Gln Ala His Cys
 275 280 285
 Asn Leu Ser Ser Thr Lys Trp Asn Asn Thr Leu Lys Gln Ile Val Thr
 290 295 300

Lys Leu Arg Glu His Phe Asn Lys Thr Ile Val Phe Asn His Ser Ser
 305 310 315 320
 5 Gly Gly Asp Pro Glu Ile Val Met His Ser Phe Asn Cys Gly Gly Glu
 325 330 335
 Phe Phe Tyr Cys Asn Thr Thr Pro Leu Phe Asn Ser Thr Trp Asn Tyr
 340 345 350
 10 Thr Tyr Thr Trp Asn Asn Thr Glu Gly Ser Asn Asp Thr Gly Arg Asn
 355 360 365
 Ile Thr Leu Gln Cys Arg Ile Lys Gln Ile Ile Asn Met Trp Gln Glu
 370 375 380
 15 Val Gly Lys Ala Met Tyr Ala Pro Pro Ile Arg Gly Gln Ile Arg Cys
 385 390 395 400
 Ser Ser Asn Ile Thr Gly Leu Leu Leu Thr Arg Asp Gly Gly Asn Asn
 405 410 415
 20 Ser Glu Thr Glu Ile Phe Arg Pro Gly Gly Gly Asp Met Arg Asp Asn
 420 425 430
 25 Trp Arg Ser Glu Leu Tyr Lys Tyr Lys Val Val Lys Ile Glu Pro Leu
 435 440 445
 Gly Val Ala Pro Thr Lys Ala *
 450 455
 30

(2) INFORMATION FOR SEQ ID NO:34:

(i) SEQUENCE CHARACTERISTICS:

- 35 (A) LENGTH: 37 base pairs
 (B) TYPE: Nucleic Acid
 (C) STRANDEDNESS: Single
 (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:34:

40 GGGAATTCGG ATCCAGAGCA GAAGACAGTG GCAATGA 37

(2) INFORMATION FOR SEQ ID NO:35:

(i) SEQUENCE CHARACTERISTICS:

- 45 (A) LENGTH: 33 base pairs
 (B) TYPE: Nucleic Acid
 (C) STRANDEDNESS: Single
 (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:35:

50 CTCGAGCTCC TGAAGACAGT CAGACTCATC AAG 33

(2) INFORMATION FOR SEQ ID NO:36:

(i) SEQUENCE CHARACTERISTICS:

- 55 (A) LENGTH: 39 base pairs
 (B) TYPE: Nucleic Acid
 (C) STRANDEDNESS: Single
 (D) TOPOLOGY: Linear

(xi) SEQUENCE DESCRIPTION: SEQ ID NO:36:

60 GGTCTAGAAG CTTTAGCCCA TAGTGCTTCC TGCTGCTCC 39

(2) INFORMATION FOR SEQ ID NO:37:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 36 base pairs
(B) TYPE: Nucleic Acid
(C) STRANDEDNESS: Single
(D) TOPOLOGY: Linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:37:
GGGCGGATCC TCGAGGTACC TGTRTGAAA GAAGCA 36

(2) INFORMATION FOR SEQ ID NO:38:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 38 base pairs
(B) TYPE: Nucleic Acid
(C) STRANDEDNESS: Single
(D) TOPOLOGY: Linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:38:
GGTCTAGAAG CTTTATGCTC CYAAGAACCC AAGGAACA 38

(2) INFORMATION FOR SEQ ID NO:39:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 7 amino acids
(B) TYPE: Amino Acid
(C) STRANDEDNESS: Single
(D) TOPOLOGY: Linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:39:
Ile Gly Pro Gly Arg Ala Phe
1 5

(2) INFORMATION FOR SEQ ID NO:40:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 7 amino acids
(B) TYPE: Amino Acid
(C) STRANDEDNESS: Single
(D) TOPOLOGY: Linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:40:
Ile Gly Pro Gly Arg Ala Trp
1 5

(2) INFORMATION FOR SEQ ID NO:41:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 7 amino acids
(B) TYPE: Amino Acid
(C) STRANDEDNESS: Single
(D) TOPOLOGY: Linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:41:
Leu Gly Pro Gly Ser Thr Phe
1 5

(2) INFORMATION FOR SEQ ID NO:42:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 7 amino acids
(B) TYPE: Amino Acid
(C) STRANDEDNESS: Single
(D) TOPOLOGY: Linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:42:

Ile Gly Pro Gly Arg Val Leu
1 5

- 5 (2) INFORMATION FOR SEQ ID NO:43:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 7 amino acids
(B) TYPE: Amino Acid
(C) STRANDEDNESS: Single
(D) TOPOLOGY: Linear
10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO:43:

Ile Gly Pro Gly Ser Ala Phe
1 5

- 15 (2) INFORMATION FOR SEQ ID NO:44:
(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 5 amino acids
(B) TYPE: Amino Acid
(C) STRANDEDNESS: Single
20 (D) TOPOLOGY: Linear
(xi) SEQUENCE DESCRIPTION: SEQ ID NO:44:

Ile Gly Pro Gly Arg
1 5

25

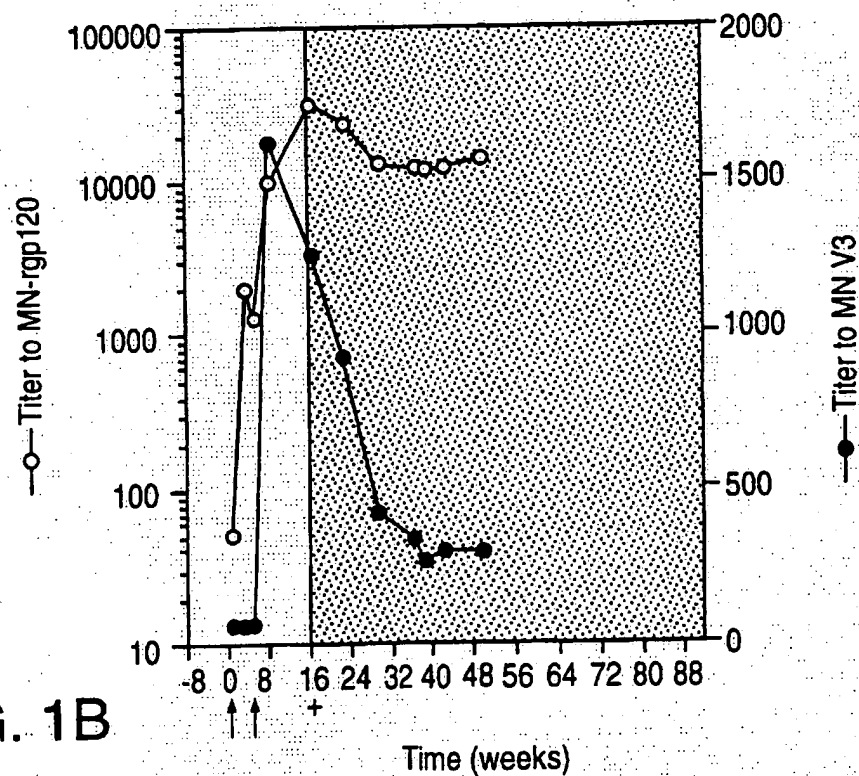
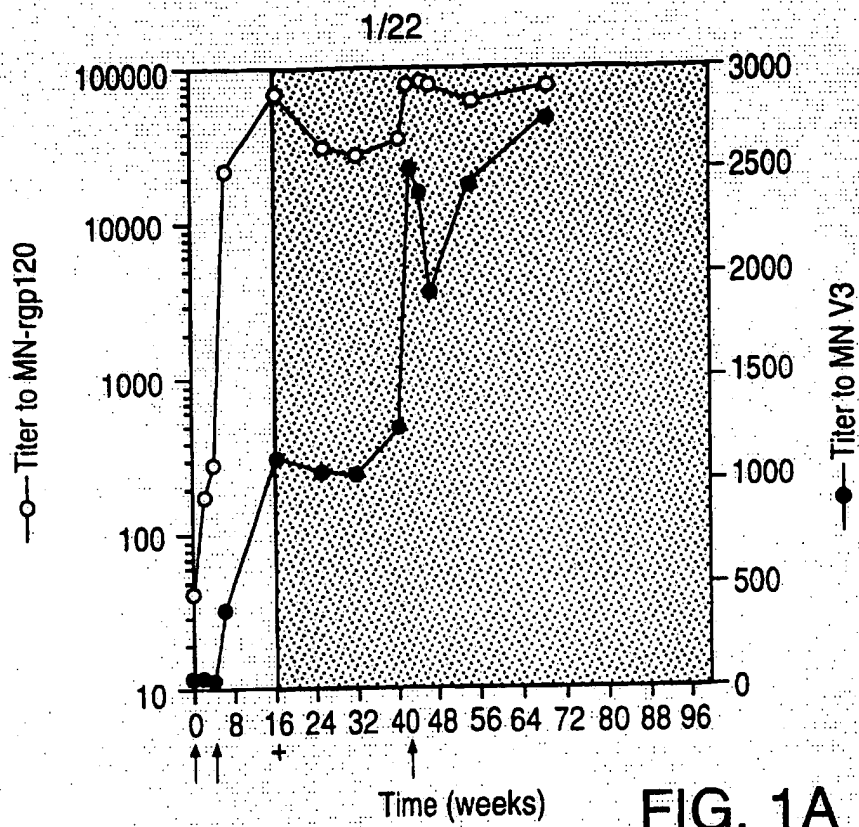
WHAT IS CLAIMED IS:

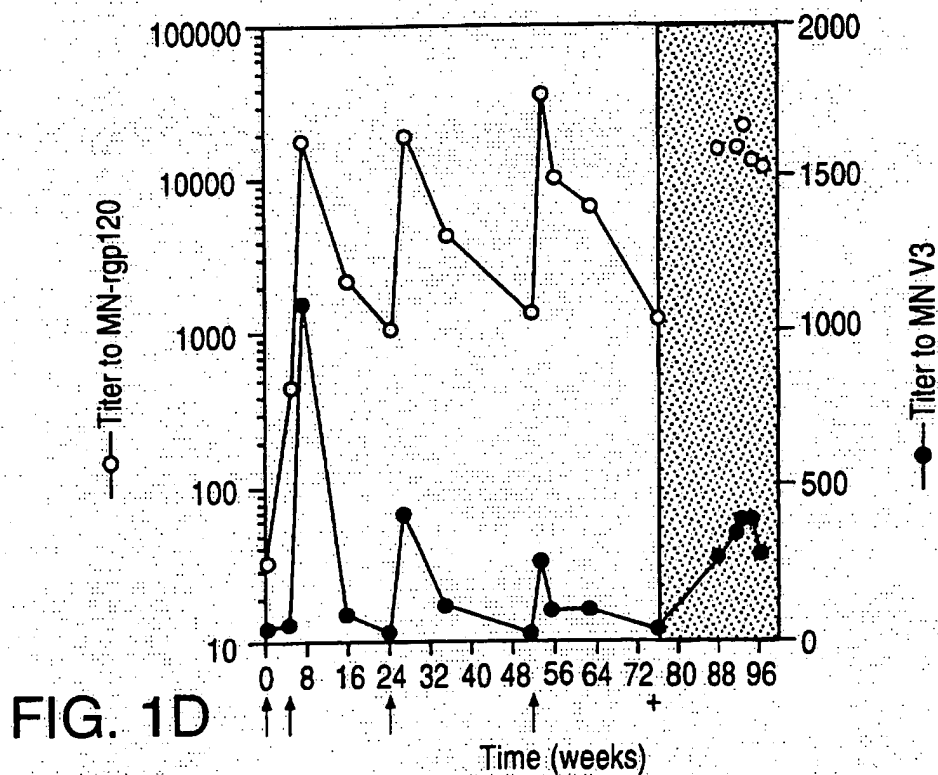
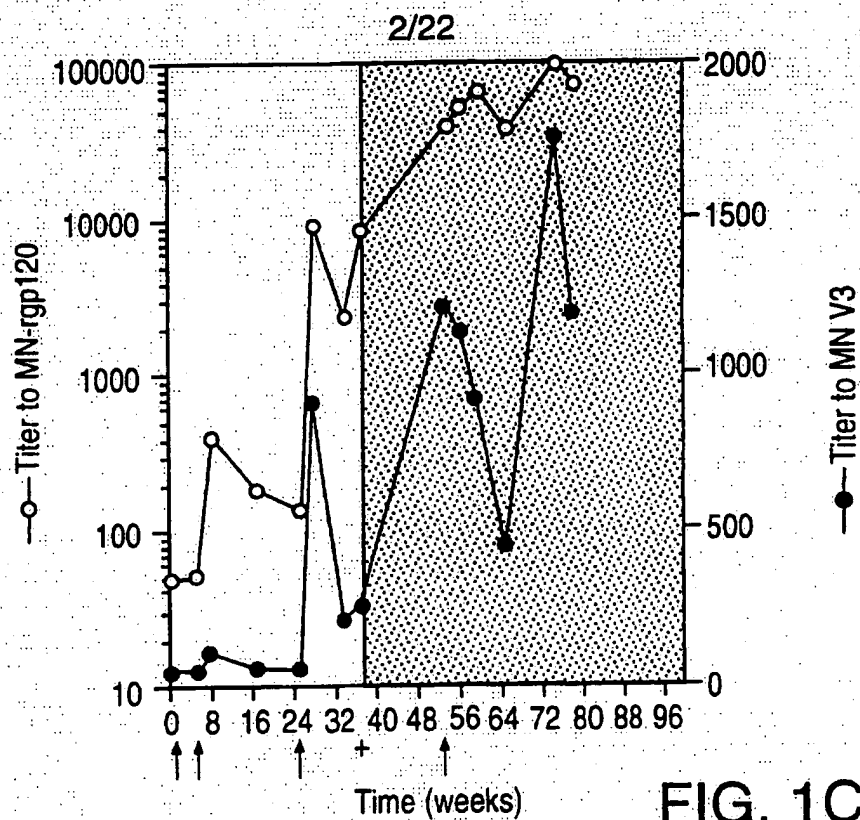
1. An isolated polypeptide comprising an HIV gp120 amino acid sequence selected from the group consisting of Sequence ID Nos. 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, and 28, and fragments thereof.
2. The polypeptide of Claim 1 wherein the polypeptide additionally comprises a flag epitope sequence.
3. The polypeptide of Claim 2 wherein the flag epitope sequence is HSV gD-1 flag epitope sequence.
4. The polypeptide of Claim 2 wherein the flag epitope sequence is fused to the HIV gp120 amino acid sequence.
5. An oligonucleotide of not more than five kilobases encoding an HIV gp120 polypeptide sequence comprising an amino acid sequence selected from the group consisting of Sequence ID Nos. 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, and 28, and fragments thereof.
6. The oligonucleotide of Claim 5 wherein the oligonucleotide includes a nucleotide sequence selected from the group consisting of Sequence ID Nos. 1, 3, 5, 7, 9, 11, 13, 15, 17, 19, 21, 23, 25, and 27, and fragments thereof.
7. The oligonucleotide of Claim 5 wherein the amino acid sequence encoded by the oligonucleotide additionally comprises a flag epitope.

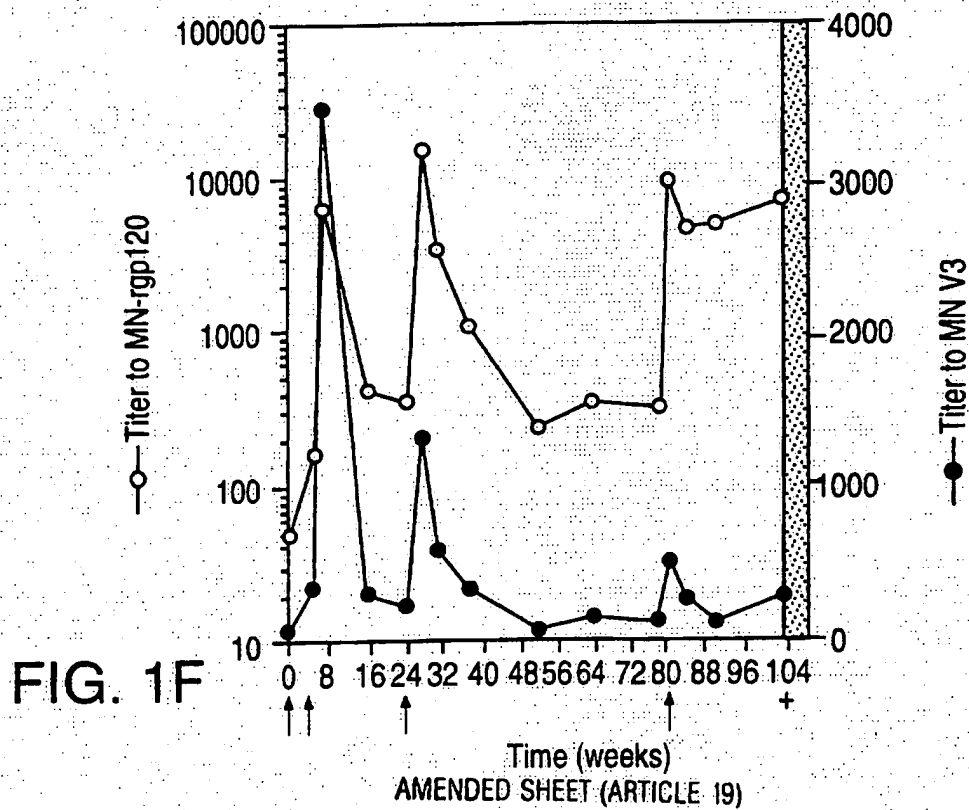
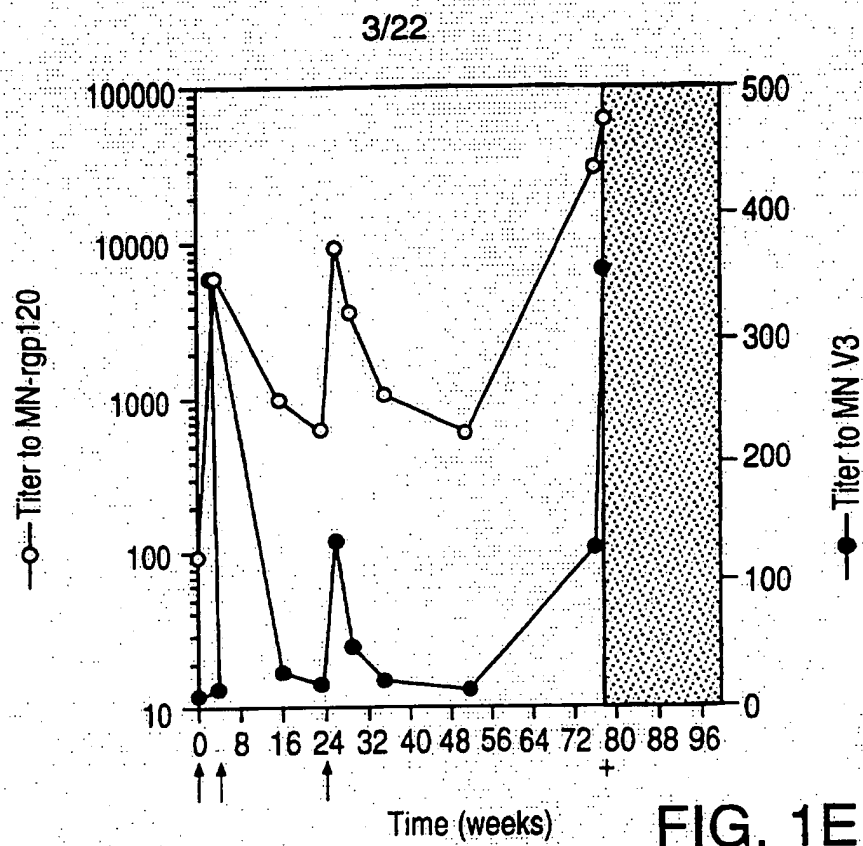
8. The oligonucleotide of Claim 5 wherein the flag epitope is HSV gD-1 flag epitope.
9. The oligonucleotide of Claim 7 wherein the flag epitope is fused to the HIV gp120 amino acid sequence.
10. A vaccine comprising gp120 MN and an HIV gp120 polypeptide sequence selected from the group consisting of Sequence ID Nos. 2, 4, 6, 8, 10, 12, 14, 16, 18, 20, 22, 24, 26, and 28, and fragments thereof in a suitable carrier.
11. A vaccine comprising:
- 15 a. a first gp120 polypeptide sequence or a fragment thereof; and
- b. a breakthrough isolate HIV gp120 polypeptide sequence or a fragment thereof from a vaccinee vaccinated with said first HIV gp120 polypeptide sequence;
- 20 wherein said HIV gp120 polypeptide sequences are in a suitable carrier.
12. The vaccine of Claim 11 wherein said first HIV gp120 polypeptide sequence comprises gp120 MN, gp120 A244, gp120 MN-GNE6 (Sequence ID No. 31), or gp120 MN-GNE8 (Sequence ID No. 33).
13. The vaccine of Claim 12 wherein said vaccine additionally comprises a second gp120 polypeptide sequence comprising gp120 MN, gp120 A244, gp120 MN-GNE6 (Sequence ID No. 31), or gp120 MN-GNE8 (Sequence ID No. 33), or a fragment thereof, wherein said second HIV gp120 polypeptide sequence is different from said first HIV gp120 polypeptide sequence.

14. The vaccine of Claim 13 wherein said first gp120 polypeptide sequence comprises gp120 MN and said second gp120 polypeptide sequence comprises gp120 A244.
- 5 15. The vaccine of Claim 14 wherein said breakthrough isolate comprises an HIV gp120 polypeptide sequence selected from the group consisting of Sequence ID Nos. 2, 4, 6, 8, 10, 12, 14, 16, 18, 10 20, 22, 24, 26, and 28, and fragments thereof in a suitable carrier.
16. The vaccine of Claim 13 wherein said first gp120 polypeptide sequence comprises gp120 MN and said second gp120 polypeptide sequence comprises gp120 MN-GNE8 (Sequence ID No. 33).
- 15 17. The vaccine of Claim 16 wherein said breakthrough isolate HIV gp120 polypeptide sequence is an HIV gp120 polypeptide sequence selected from the group consisting of Sequence ID Nos. 2, 4, 6, 8, 10, 12, 20 14, 16, 18, 20, 22, 24, 26, and 28, and fragments thereof, in a suitable carrier.
- 25 18. The vaccine of Claim 13 wherein said breakthrough isolate HIV gp120 polypeptide is from a vaccinee vaccinated with said first and second HIV gp120 polypeptide sequences.

19. A method for making an HIV vaccine comprising adding an HIV gp120 polypeptide sequence or fragments thereof from a breakthrough isolate from a vaccinee to the vaccine with which the vaccinee was vaccinated.
20. The vaccine of Claim 11 wherein said first gp120 polypeptide sequence is from a macrophage-tropic HIV-1 strain.
21. The vaccine of Claim 11 wherein said first gp120 polypeptide sequence is from a T-cell-tropic HIV-1 strain.
22. The vaccine of Claim 21 wherein said vaccine additionally comprises a second gp120 polypeptide sequence or a fragment, from a macrophage-tropic HIV-1 strain.
23. The vaccine of Claim 22 wherein said first and second gp120 polypeptide sequences bind to different chemokine receptors.
24. The vaccine of Claim 23 wherein said first gp120 polypeptide sequence binds to CC-CKR-5 and said second gp 120 polypeptide sequence binds to CXC-CKR-4.
25. The vaccine of Claim 11 wherein said vaccine additionally comprises an virus engineered to induce a cytotoxic T-cell response.







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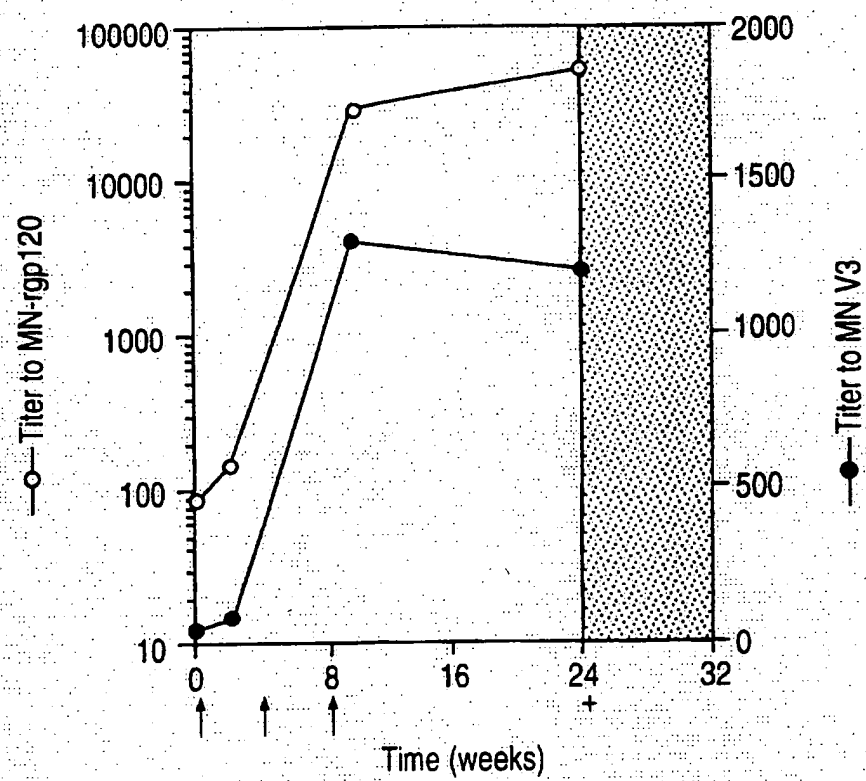


FIG. 1G

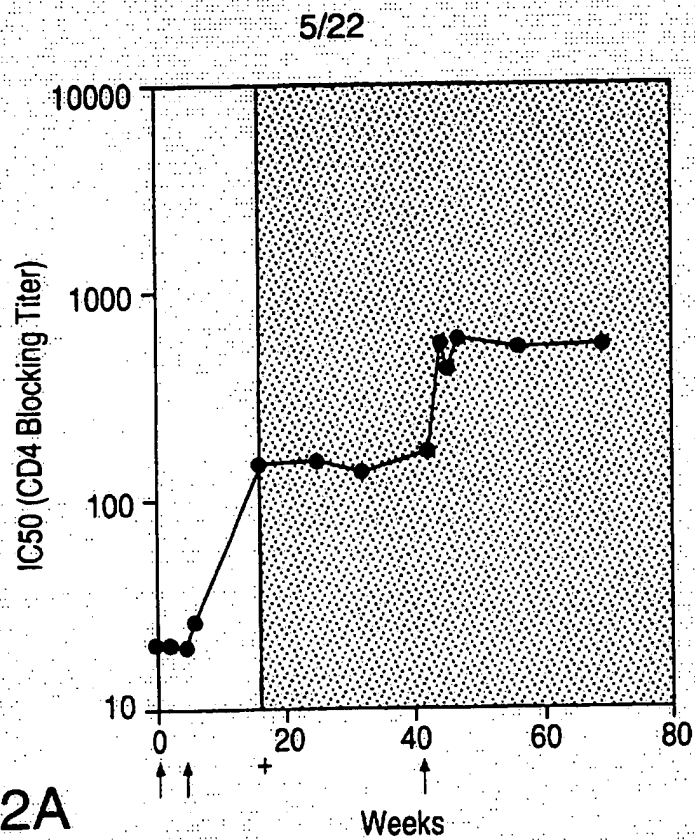


FIG. 2A

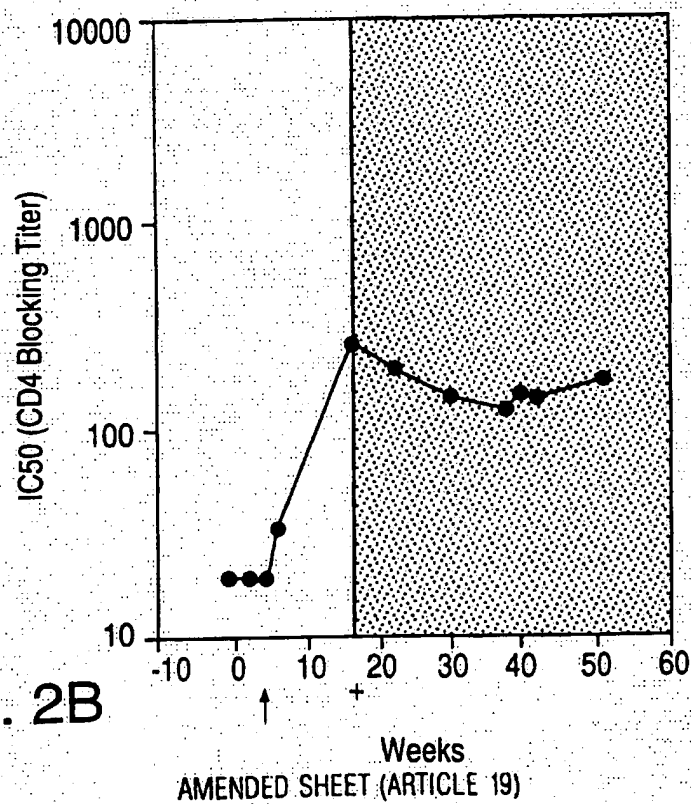


FIG. 2B

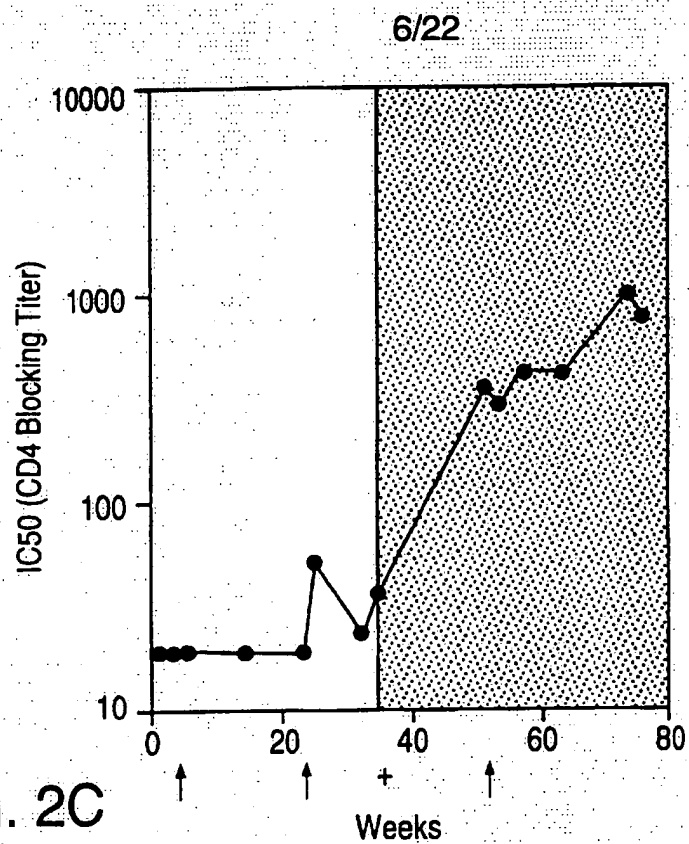


FIG. 2C

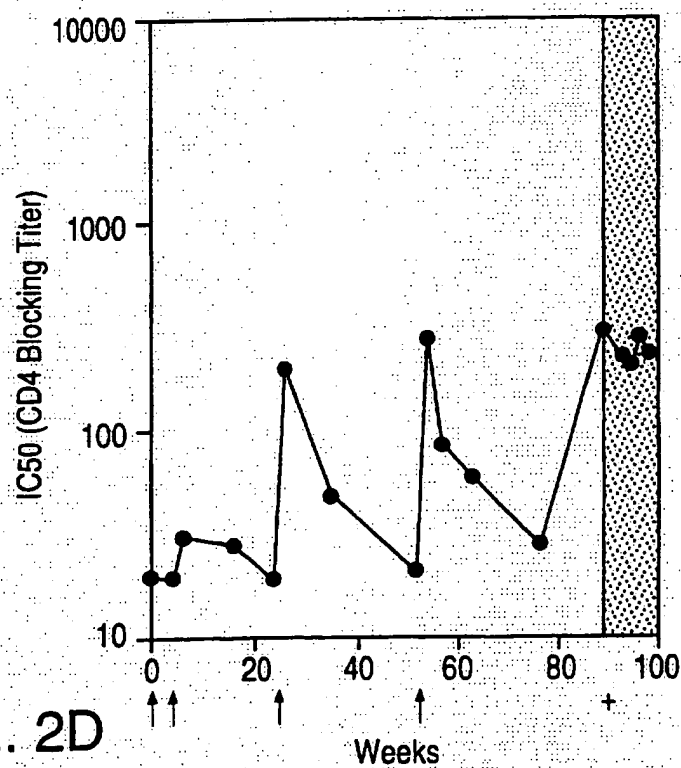
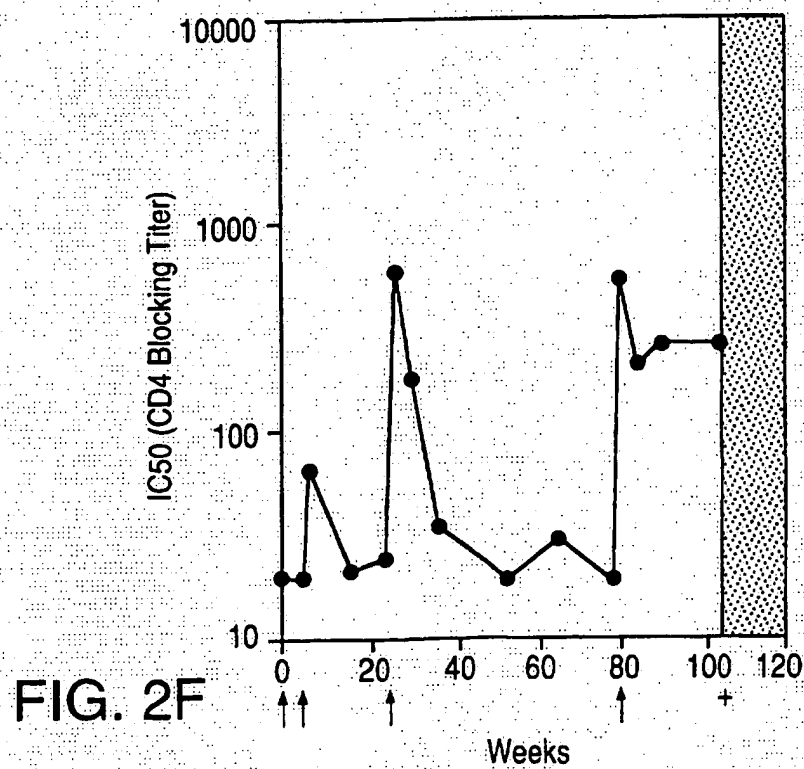
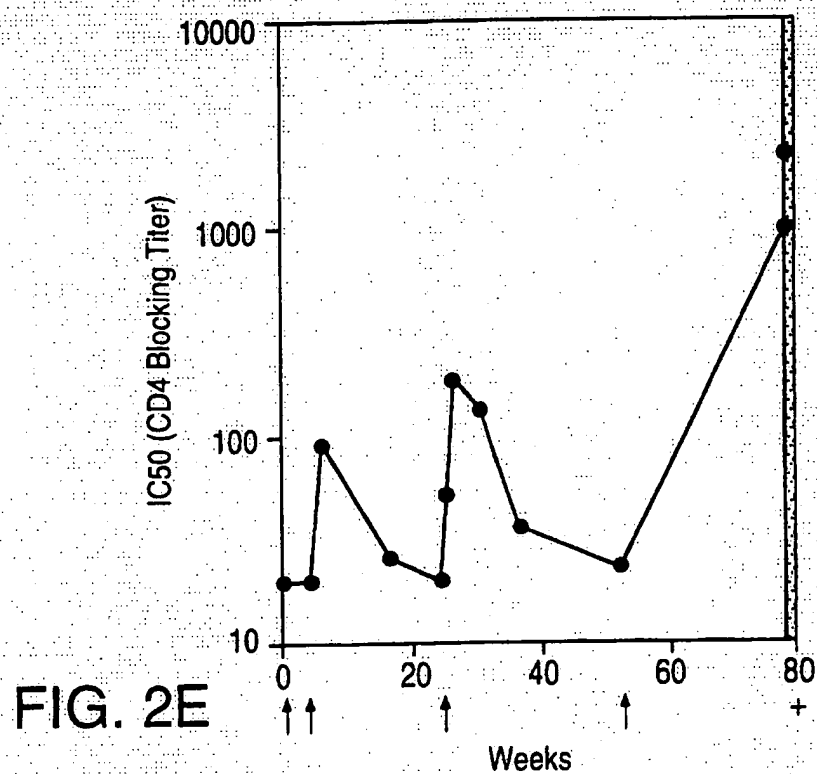


FIG. 2D

AMENDED SHEET (ARTICLE 19)

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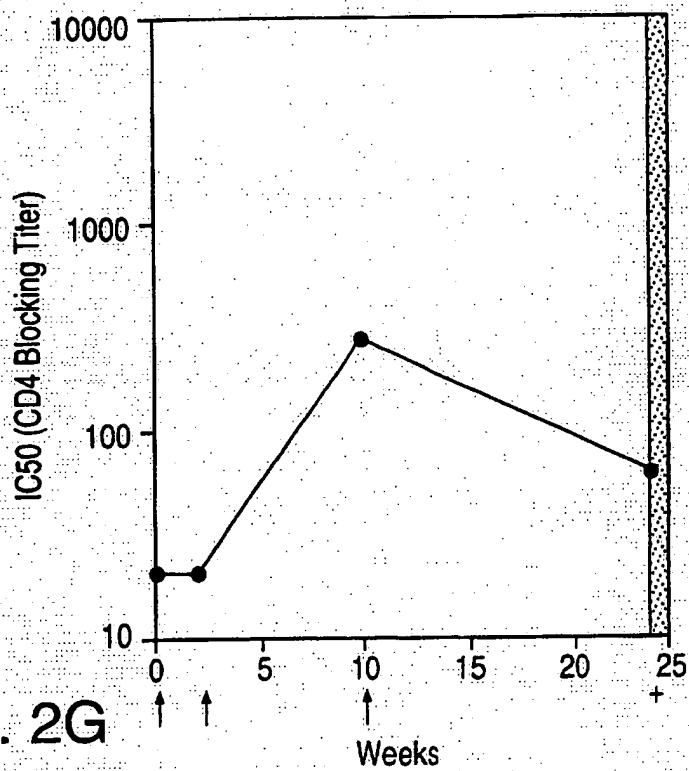


FIG. 2G

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| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
|-------|----|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
| C6.1 | 41 | V | P | V | W | K | E | A | T | T | T | L | F | C | A | S | D | A | K | A | Y | D | T | E | V | H | N | V | W | A | T | H | A | C | V | P | T | D | P | N | P | Q | E | E | M | V | L | E | E |
| C6.5 | 41 | V | P | V | W | K | E | A | T | T | T | L | F | C | A | S | D | A | K | A | Y | D | T | E | V | H | N | V | W | A | T | H | A | C | V | P | T | D | P | N | P | Q | E | E | M | V | L | E | E |
| C8.3 | 41 | V | P | V | W | K | E | A | T | T | T | L | F | C | A | S | D | A | K | A | Y | D | T | E | V | H | N | V | W | A | T | H | A | C | V | P | T | D | P | N | P | Q | E | E | M | V | L | E | E |
| C8.6 | 41 | V | P | V | W | K | E | A | T | T | T | L | F | C | A | S | D | A | K | A | Y | D | T | E | V | H | N | V | W | A | T | H | A | C | V | P | T | D | P | N | P | Q | E | E | M | V | L | E | E |
| C15.2 | 41 | V | P | V | W | K | E | A | T | T | T | L | F | C | A | S | D | A | K | A | Y | D | T | E | V | H | N | V | W | A | T | H | A | C | V | P | T | D | P | N | P | Q | E | E | M | V | L | E | E |
| C15.3 | 41 | V | P | V | W | K | E | A | T | T | T | L | F | C | A | S | D | A | K | A | Y | D | T | E | V | H | N | V | W | A | T | H | A | C | V | P | T | D | P | N | P | Q | E | E | M | V | L | E | E |
| C7.2 | 41 | V | P | V | W | K | E | A | T | T | T | L | F | C | A | S | D | A | K | A | Y | D | T | E | V | H | N | V | W | A | T | H | A | C | V | P | T | D | P | N | P | Q | E | E | M | V | L | E | E |
| C7.10 | 41 | V | P | V | W | K | E | A | T | T | T | L | F | C | A | S | D | A | K | A | Y | D | T | E | V | H | N | V | W | A | T | H | A | C | V | P | T | D | P | N | P | Q | E | E | M | V | L | E | E |
| C11.5 | 41 | V | P | V | W | K | E | A | T | T | T | L | F | C | A | S | D | A | K | A | Y | D | T | E | V | H | N | V | W | A | T | H | A | C | V | P | T | D | P | N | P | Q | E | E | M | V | L | E | E |
| C11.7 | 41 | V | P | V | W | K | E | A | T | T | T | L | F | C | A | S | D | A | K | A | Y | D | T | E | V | H | N | V | W | A | T | H | A | C | V | P | T | D | P | N | P | Q | E | E | M | V | L | E | E |
| C10.5 | 41 | V | P | V | W | K | E | A | T | T | T | L | F | C | A | S | D | A | K | A | Y | D | T | E | V | H | N | V | W | A | T | H | A | C | V | P | T | D | P | N | P | Q | E | E | M | V | L | E | E |
| C10.7 | 41 | V | P | V | W | K | E | A | T | T | T | L | F | C | A | S | D | A | K | A | Y | D | T | E | V | H | N | V | W | A | T | H | A | C | V | P | T | D | P | N | P | Q | E | E | M | V | L | E | E |
| C17.1 | 41 | V | P | V | W | K | E | A | T | T | T | L | F | C | A | S | D | A | K | A | Y | D | T | E | V | H | N | V | W | A | T | H | A | C | V | P | T | D | P | N | P | Q | E | E | M | V | | | |

FIG. 3A

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| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
|-------|-----|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
| C6.1 | 240 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | A | H | G | I | K | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| C6.5 | 240 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | K | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| C8.3 | 223 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | R | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| C8.6 | 223 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | R | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| C15.2 | 225 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | R | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| C15.3 | 225 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | R | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| C7.2 | 223 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | R | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| C7.10 | 223 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | R | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| C11.5 | 236 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | R | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| C11.7 | 236 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | R | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| C10.5 | 224 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | R | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| C10.7 | 224 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | R | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| C17.1 | 214 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | K | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| C17.3 | 214 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | K | P | V | V | S | T | Q | L | L | N | G | S | L | A |
| MNGNE | 226 | A | G | F | A | I | L | K | C | R | D | K | F | N | G | T | G | P | C | K | N | V | S | T | V | Q | C | T | H | G | I | R | P | V | V | S | T | Q | L | L | N | G | S | L | A |

FIG. 3E

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| | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
|-------|-----|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|---|
| C6.1 | 340 | I | R | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| C6.5 | 340 | I | R | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| C8.3 | 323 | I | I | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| C8.6 | 323 | I | I | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| C15.2 | 325 | I | I | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| C15.3 | 325 | I | I | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| C7.2 | 323 | I | V | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| C7.10 | 323 | I | V | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| C11.5 | 336 | I | I | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| C11.7 | 336 | I | I | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| C10.5 | 324 | I | I | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| C10.7 | 324 | I | I | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| C17.1 | 314 | I | I | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| C17.3 | 314 | I | I | G | D | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |
| MNGNE | 326 | I | K | G | T | I | R | Q | A | H | C | N | I | S | G | A | K | W | N | N | T | L | L | K | K | V | V | K | L | K | E | E | Q | Q | F | F | K | K | R | Q | F | G | P | N | K | T | I | V | F | N | F | S |

FIG. 3G

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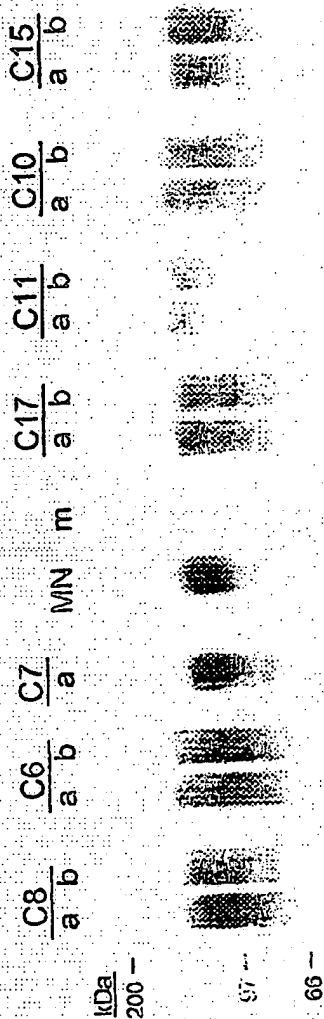


FIG. 4

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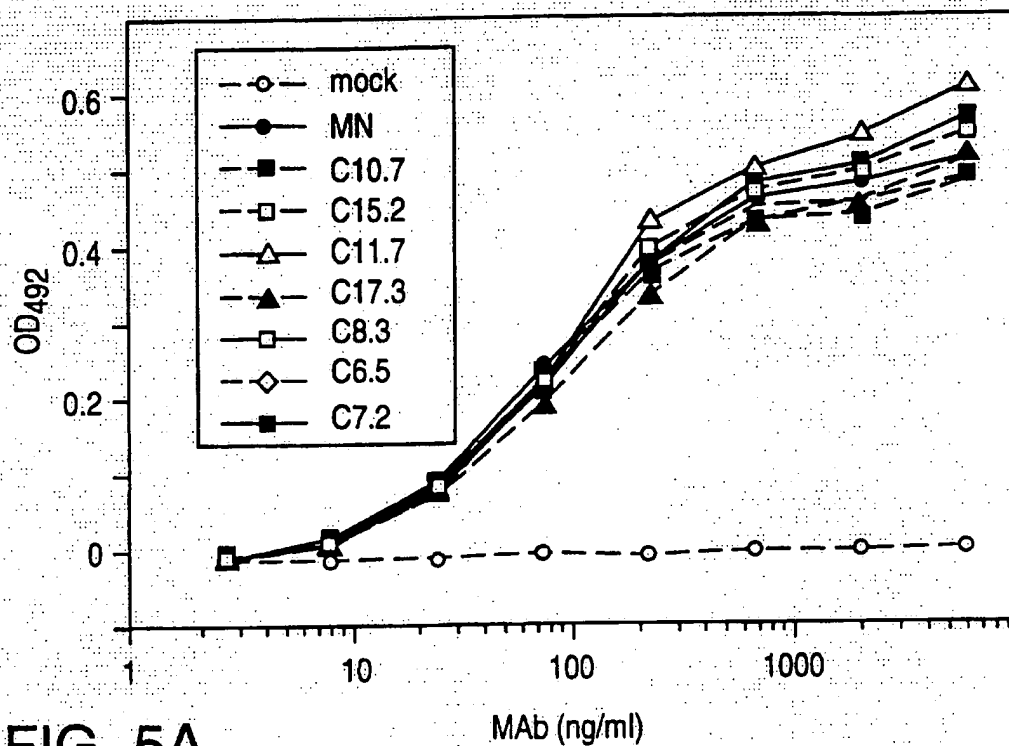


FIG. 5A

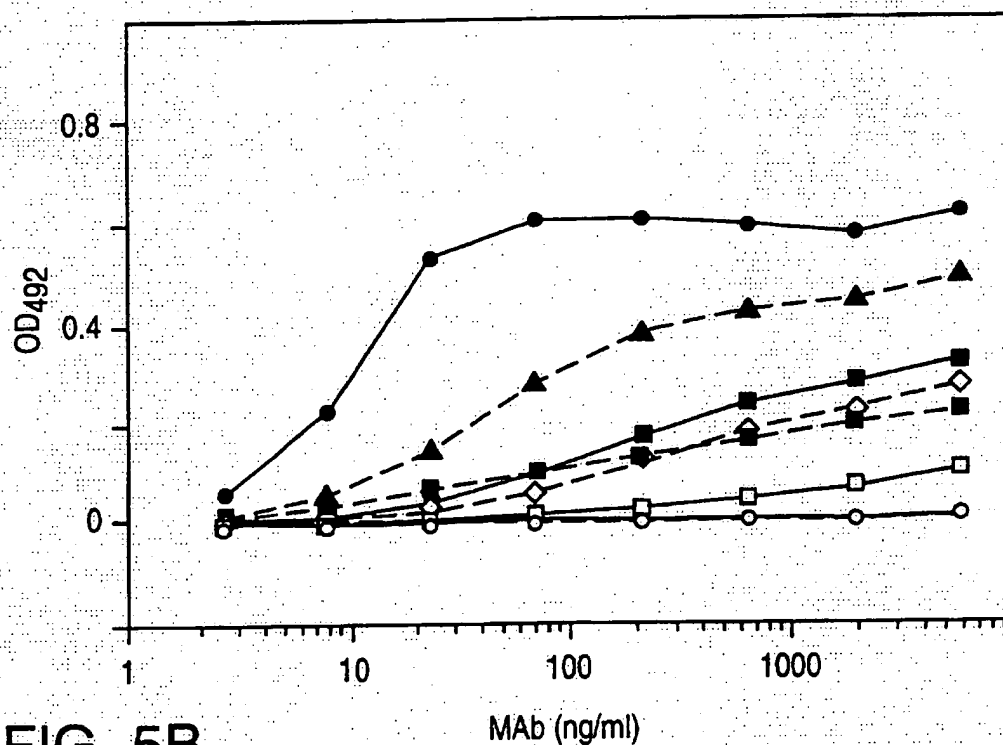


FIG. 5B

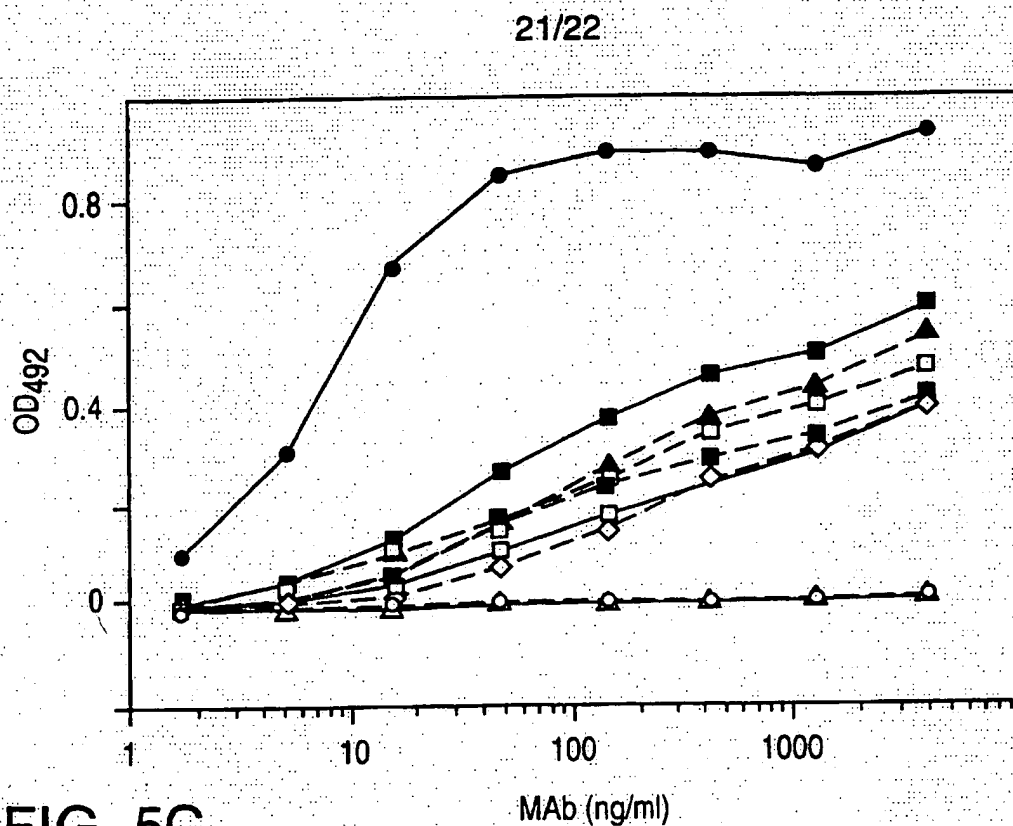


FIG. 5C

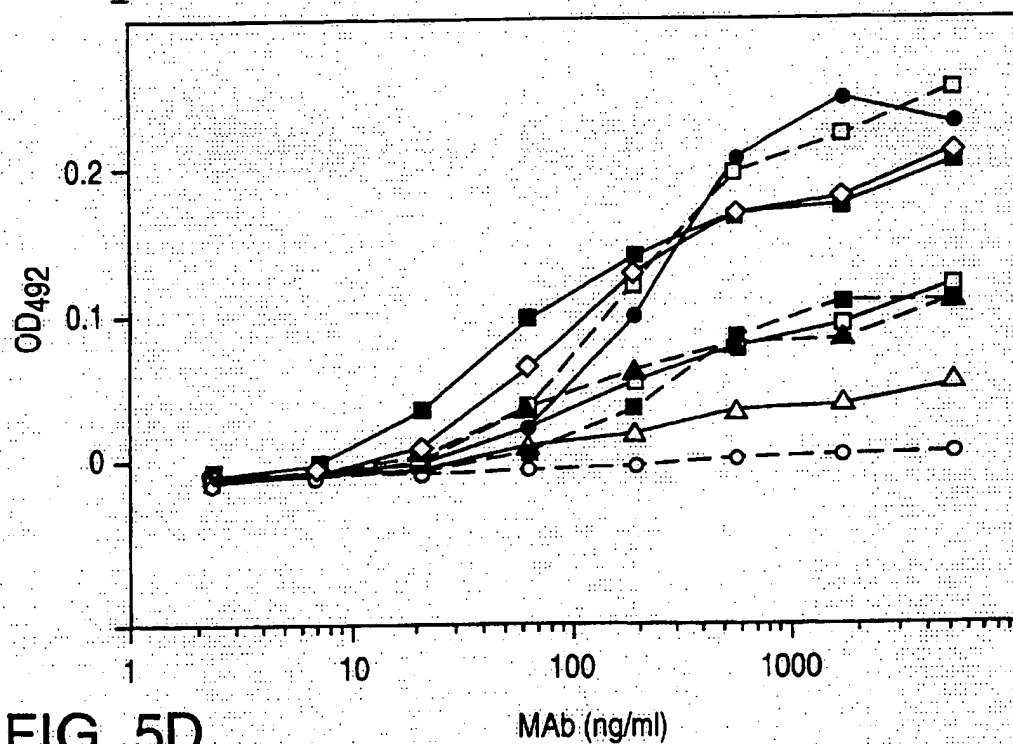


FIG. 5D

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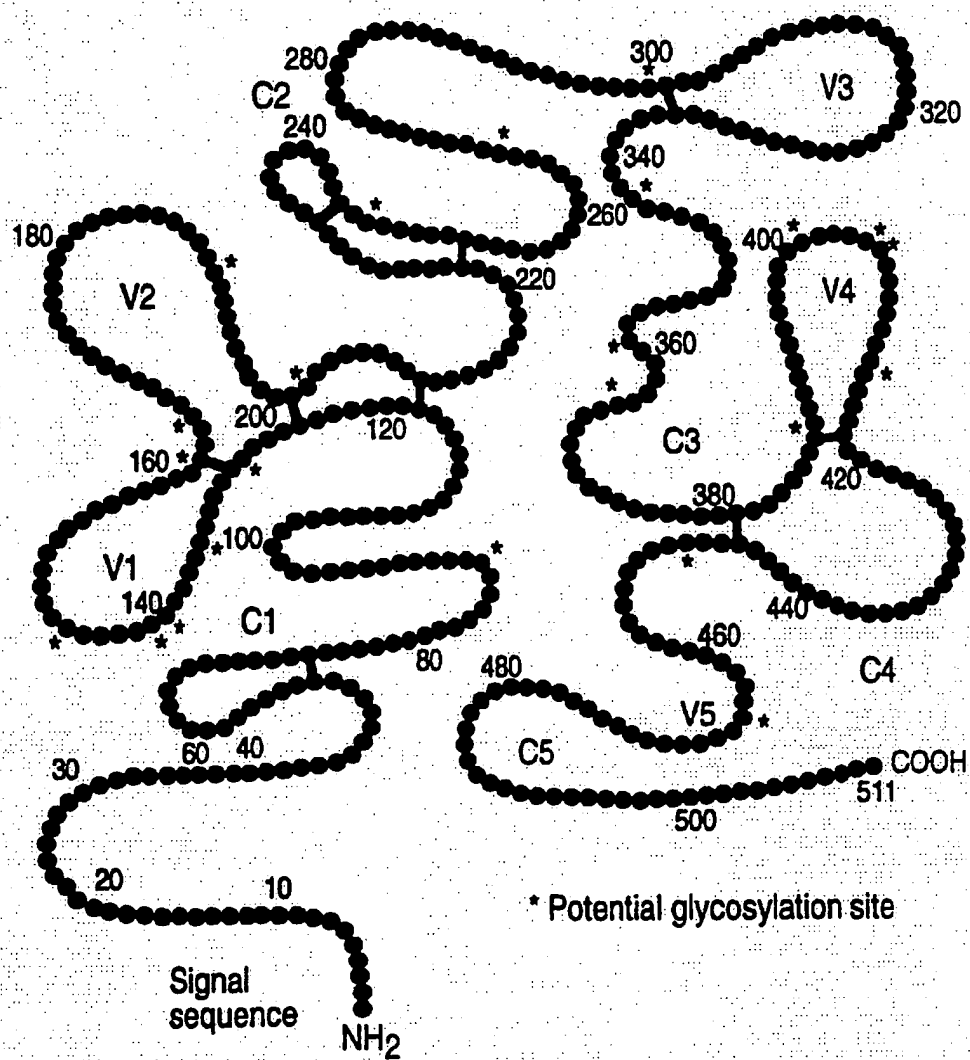


FIG. 6

INTERNATIONAL SEARCH REPORT

International Application No.
PCT/US 97/09690

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C12N15/49 C07K14/16 A61K39/21

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C12N C07K A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

| Category * | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|------------|--|-----------------------|
| X | WO 94 28929 A (GENETECH, INC.) 22 December 1994 see page 56 SEQ. ID. NO. 25. see page 50, line 14 - line 31 --- | 1-18, 20-25 |
| X | P.W. BERMAN ET AL.: "Genetic and immunologic characterization of viruses infecting MN-rgp120 vaccinated volunteers" ONE WORLD, ONE HOPE: XI INTERNATIONAL CONFERENCE ON AIDS, vol. 10, no. supplement 3, 7 - 12 July 1996, VANCOUVER, CANADA, page 10 XP002045307 See "Methods" in Abstract Mo.A.285 --- -/-- | 1,5,6 |



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

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Date of the actual completion of the international search

30 October 1997

Date of mailing of the international search report

26 -11- 1997

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

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|------------|---|-----------------------|
| A | M.J. MCEL RATH ET AL.: "Human immunodeficiency virus type 1 infection despite prior immunization with a recombinant envelope vaccine regimen" PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF USA, vol. 93, no. 9, 30 April 1996, WASHINGTON US, pages 3972-3977, XP002045308 see page 3976, last paragraph; figure 1 --- | 19 |
| T | P.W. BERMAN ET AL.: "Genetic and immunologic characterization of viruses infecting MN-rgp120-vaccinated volunteers" THE JOURNAL OF INFECTIOUS DISEASES, vol. 176, no. 2, August 1997, pages 384-397, XP002045309 see the whole document ----- | 1-25 |

information on patent family members

PCT/US 97/09690

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